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REQUEST FOR FILING A CONTINUATION PATENT APPLICATION UNDER 37 CFR 1.53(b)

DOCKET NUMBER	ANTICIPATED CLASSIFICATION OF THIS APPLICATION		PRIOR APPLICATION SERIAL NO.	PRIOR APPLICATION EXAMINER	ART UNIT
JHV-010.03	CLASS	SUBCLASS	09/685,244	Not Assigned	Not Assigned

Address to: Box Patent Application  
Assistant Commissioner for Patents  
Washington, D.C. 20231

This is a request for filing a (X) continuation ( ) divisional application under 37 CFR 1.53(b), of pending prior application Number 09/685,244, filed on October 10, 2000 entitled "Inhibitors of Hedgehog Signaling Pathways, Compositions and Uses Related Thereto."

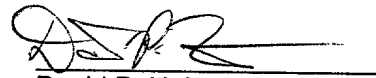
1. I hereby verify that the papers are a true copy of the latest signed prior application number 09/685,244, and further that all statements made herein of my own knowledge are true; and further that these statements were made with the knowledge that willful false statements and the like are made punishable by fine or imprisonment, or both, under section 1001 of Title 18 of the United States Code and that such willful statements may jeopardize the validity of the application or any patent issuing thereon.

CLAIM	(1) FOR	(2) NUMBER FILED	(3) NUMBER EXTRA	(4) RATE	(5) CALCULATIONS
	TOTAL CLAIMS (37 CFR 1.16(c))			x \$18.00=	\$
	INDEPENDENT CLAIMS (37 CFR 1.16(b))		0	x \$78.00=	\$
	MULTIPLE DEPENDENT CLAIMS (if applicable) (37 CFR 1.16(d))			+ \$260.00=	\$
				BASIC FEE (37 CFR 1.16(a))	\$
				Total of above Calculations =	\$
	*Reduction by 50% for filing by small entity (Note 37 CFR 1.9, 1.27, 1.28)				-\$
	TOTAL=				\$

2. ( ) A verified statement to establish small entity status under 37 CFR 1.9 and 1.27  
( ) is enclosed.  
( ) was filed in prior application and such status is still proper and desired (37 CFR 1.28(a)).
3. ( ) The Commissioner is hereby authorized to charge any fees which may be required under 37 CFR 1.16 and 1.17, or credit any overpayment to Deposit Account No. 06-1448. A duplicate copy of this sheet is enclosed.
- (X) Filing fees are not being paid at this time.
- ( ) Cancel in this application original claims and of the prior application and enter the Preliminary Amendment attached hereto before calculating the filing fee. (At least one original independent claim must be retained for filing purposes.)

6. (X) The inventor(s) of the invention being claimed in this application is (are): Philip A. Beachy, Michael K. Cooper, and Jeffrey A. Porter.
7. ( ) This application is being filed by less than all the inventors named in the prior application. In accordance with 37 CFR 1.60(b), the Commissioner is requested to delete the name(s) of the following person or persons who are not inventors of the invention being claimed in this application:
8. (X) Amend the specification by inserting on page 1, line 4, after "This application is", --a continuation of U.S. Application No. 09/685,244 filed October 10, 2000, which is--
9. ( ) Transfer the drawings from the pending prior application to this application and abandon said prior application as of the filing date accorded this application. A duplicate copy of this sheet is enclosed for filing in prior application. (May only be used if signed by person authorized by 37 CFR 1.138 and before payment of issue fee.)
10. ( ) Diskette with Sequence Listing.
11. ( ) Priority of foreign application number \_\_\_\_\_, filed on \_\_\_\_\_, in \_\_\_\_\_ is claimed under 35 U.S.C 119.
- ( ) The certified copy has been filed in prior application number \_\_\_\_/\_\_\_\_\_, filed \_\_\_\_\_.
12. ( ) A preliminary amendment is enclosed.
13. ( ) The prior application is assigned of record to: \_\_\_\_\_.
14. (X) Also enclosed: Copy of unexecuted Declaration from application number 09/685,244.
15. ( ) The power of attorney in the prior application is to: \_\_\_\_\_.
- a. ( ) The power of attorney appears in the prior application.
- b. ( ) Since the power does not appear in the original paper, a copy of the power in the prior application is enclosed.
- c. (X) Address all future correspondence to: (May only be completed by applicant, or attorney or agent of record.)
- Patent Group
- Foley, Hoag & Eliot LLP
- One Post Office Square
- Boston, Massachusetts 02109-2170

November 8, 2000  
Date

  
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Date of Signature  
and of Mail Deposit

  
Robert King

20001108 14250000

*Inhibitors of Hedgehog Signaling Pathways,  
Compositions and Uses Related Thereto*

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This application is a continuation-in-part of PCT Application No. US99/07811,  
5 filed April 9, 1999, based on U.S. Application No. 09/090,622, filed June 4, 1998, which  
claims priority to U.S. Provisional Application Nos. 60/081,186 and 60/081,263, filed  
April 9, 1998, the specifications of all of which are hereby incorporated herein by  
reference in their entirety.

10 **Background of the Invention**

Pattern formation is the activity by which embryonic cells form ordered spatial  
arrangements of differentiated tissues. The physical complexity of higher organisms arises  
during embryogenesis through the interplay of cell-intrinsic lineage and cell-extrinsic  
signaling. Inductive interactions are essential to embryonic patterning in vertebrate  
15 development from the earliest establishment of the body plan, to the patterning of the  
organ systems, to the generation of diverse cell types during tissue differentiation  
(Davidson, E., (1990) Development 108: 365-389; Gurdon, J. B., (1992) Cell 68: 185-199;  
Jessell, T. M. et al., (1992) Cell 68: 257-270). The effects of developmental cell  
interactions are varied. Typically, responding cells are diverted from one route of cell  
20 differentiation to another by inducing cells that differ from both the uninduced and  
induced states of the responding cells (inductions). Sometimes cells induce their neighbors  
to differentiate like themselves (homeogenetic induction); in other cases a cell inhibits its  
neighbors from differentiating like itself. Cell interactions in early development may be  
sequential, such that an initial induction between two cell types leads to a progressive  
25 amplification of diversity. Moreover, inductive interactions occur not only in embryos, but  
in adult cells as well, and can act to establish and maintain morphogenetic patterns as well  
as induce differentiation (J.B. Gurdon (1992) Cell 68:185-199).

Members of the *Hedgehog* family of signaling molecules mediate many important  
short- and long-range patterning processes during invertebrate and vertebrate development.  
30 In the fly, a single *hedgehog* gene regulates segmental and imaginal disc patterning. In  
contrast, in vertebrates, a *hedgehog* gene family is involved in the control of left-right

asymmetry, polarity in the CNS, somites and limb, organogenesis, chondrogenesis and spermatogenesis.

The first *hedgehog* gene was identified by a genetic screen in the fruitfly *Drosophila melanogaster* (Nüsslein-Volhard, C. and Wieschaus, E. (1980) Nature 287, 795-801). This screen identified a number of mutations affecting embryonic and larval development. In 1992 and 1993, the molecular nature of the *Drosophila hedgehog* (*hh*) gene was reported (C.F., Lee et al. (1992) Cell 71, 33-50), and since then, several *hedgehog* homologues have been isolated from various vertebrate species. While only one *hedgehog* gene has been found in *Drosophila* and other invertebrates, multiple *Hedgehog* genes are present in vertebrates.

The vertebrate family of *hedgehog* genes includes at least four members, e.g., paralogs of the single drosophila *hedgehog* gene. Exemplary *hedgehog* genes and proteins are described in PCT publications WO 95/18856 and WO 96/17924. Three of these members, herein referred to as Desert *hedgehog* (*Dhh*), Sonic *hedgehog* (*Shh*) and Indian *hedgehog* (*Ihh*), apparently exist in all vertebrates, including fish, birds, and mammals. A fourth member, herein referred to as tiggie-winkle *hedgehog* (*Thh*), appears specific to fish. Desert *hedgehog* (*Dhh*) is expressed principally in the testes, both in mouse embryonic development and in the adult rodent and human; Indian *hedgehog* (*Ihh*) is involved in bone development during embryogenesis and in bone formation in the adult; and, *Shh*, which as described above, is primarily involved in morphogenic and neuroinductive activities. Given the critical inductive roles of *hedgehog* polypeptides in the development and maintenance of vertebrate organs, the identification of *hedgehog* interacting proteins is of paramount significance in both clinical and research contexts.

The various *Hedgehog* proteins consist of a signal peptide, a highly conserved N-terminal region, and a more divergent C-terminal domain. In addition to signal sequence cleavage in the secretory pathway (Lee, J.J. et al. (1992) Cell 71:33-50; Tabata, T. et al. (1992) Genes Dev. 2635-2645; Chang, D.E. et al. (1994) Development 120:3339-3353), *Hedgehog* precursor proteins undergo an internal autoproteolytic cleavage which depends on conserved sequences in the C-terminal portion (Lee et al. (1994) Science 266:1528-1537; Porter et al. (1995) Nature 374:363-366). This autocleavage leads to a 19 kD N-terminal peptide and a C-terminal peptide of 26-28 kD (Lee et al. (1992) supra; Tabata et al. (1992) supra; Chang et al. (1994) supra; Lee et al. (1994) supra; Bumcrot, D.A., et al.

(1995) Mol. Cell. Biol. 15:2294-2303; Porter *et al.* (1995) supra; Ekker, S.C. *et al.* (1995) Curr. Biol. 5:944-955; Lai, C.J. *et al.* (1995) Development 121:2349-2360). The N-terminal peptide stays tightly associated with the surface of cells in which it was synthesized, while the C-terminal peptide is freely diffusible both *in vitro* and *in vivo*

5 (Porter *et al.* (1995) Nature 374:363; Lee *et al.* (1994) supra; Bumcrot *et al.* (1995) supra; Mart', E. *et al.* (1995) Development 121:2537-2547; Roelink, H. *et al.* (1995) Cell 81:445-455). Interestingly, cell surface retention of the N-terminal peptide is dependent on autocleavage, as a truncated form of HH encoded by an RNA which terminates precisely at the normal position of internal cleavage is diffusible *in vitro* (Porter *et al.* (1995) supra)

10 and *in vivo* (Porter, J.A. *et al.* (1996) Cell 86, 21-34). Biochemical studies have shown that the autoproteolytic cleavage of the HH precursor protein proceeds through an internal thioester intermediate which subsequently is cleaved in a nucleophilic substitution. It is likely that the nucleophile is a small lipophilic molecule which becomes covalently bound to the C-terminal end of the N-peptide (Porter *et al.* (1996) supra), tethering it to the cell

15 surface. The biological implications are profound. As a result of the tethering, a high local concentration of N-terminal Hedgehog peptide is generated on the surface of the Hedgehog producing cells. It is this N-terminal peptide which is both necessary and sufficient for short- and long-range Hedgehog signaling activities in *Drosophila* and vertebrates (Porter *et al.* (1995) supra; Ekker *et al.* (1995) supra; Lai *et al.* (1995) supra; Roelink, H. *et al.*

20 (1995) Cell 81:445-455; Porter *et al.* (1996) supra; Fietz, M.J. *et al.* (1995) Curr. Biol. 5:643-651; Fan, C.-M. *et al.* (1995) Cell 81:457-465; Mart', E., *et al.* (1995) Nature 375:322-325; Lopez-Martinez *et al.* (1995) Curr. Biol. 5:791-795; Ekker, S.C. *et al.* (1995) Development 121:2337-2347; Forbes, A.J. *et al.* (1996) Development 122:1125-1135).

HH has been implicated in short- and long-range patterning processes at various

25 sites during *Drosophila* development. In the establishment of segment polarity in early embryos, it has short-range effects which appear to be directly mediated, while in the patterning of the imaginal discs, it induces long range effects via the induction of secondary signals.

In vertebrates, several *hedgehog* genes have been cloned in the past few years. Of

30 these genes, *Shh* has received most of the experimental attention, as it is expressed in different organizing centers which are the sources of signals that pattern neighboring tissues. Recent evidence indicates that *Shh* is involved in these interactions.

The expression of *Shh* starts shortly after the onset of gastrulation in the presumptive midline mesoderm, the node in the mouse (Chang *et al.* (1994) supra; Echelard, Y. *et al.* (1993) Cell 75:1417-1430), the rat (Roelink, H. *et al.* (1994) Cell 76:761-775) and the chick (Riddle, R.D. *et al.* (1993) Cell 75:1401-1416), and the shield  
 5 in the zebrafish (Ekker *et al.* (1995) supra; Krauss, S. *et al.* (1993) Cell 75:1431-1444). In chick embryos, the *Shh* expression pattern in the node develops a left-right asymmetry, which appears to be responsible for the left-right situs of the heart (Levin, M. *et al.* (1995) Cell 82:803-814).

In the CNS, *Shh* from the notochord and the floorplate appears to induce ventral  
 10 cell fates. When ectopically expressed, *Shh* leads to a ventralization of large regions of the mid- and hindbrain in mouse (Echelard *et al.* (1993) supra; Goodrich, L.V. *et al.* (1996) Genes Dev. 10:301-312), *Xenopus* (Roelink, H. *et al.* (1994) supra; Ruiz i Altaba, A. *et al.* (1995) Mol. Cell. Neurosci. 6:106-121), and zebrafish (Ekker *et al.* (1995) supra; Krauss  
 15 *et al.* (1993) supra; Hammerschmidt, M., *et al.* (1996) Genes Dev. 10:647-658). In explants of intermediate neuroectoderm at spinal cord levels, *Shh* protein induces floorplate and motor neuron development with distinct concentration thresholds, floor plate at high and motor neurons at lower concentrations (Roelink *et al.* (1995) supra; Mart'  
 20 *et al.* (1995) supra; Tanabe, Y. *et al.* (1995) Curr. Biol. 5:651-658). Moreover, antibody blocking suggests that *Shh* produced by the notochord is required for notochord-mediated induction of motor neuron fates (Mart' *et al.* (1995) supra). Thus, high concentration of *Shh* on the surface of *Shh*-producing midline cells appears to account for the contact-mediated induction of floorplate observed *in vitro* (Placzek, M. *et al.* (1993) Development  
 25 117:205-218), and the midline positioning of the floorplate immediately above the notochord *in vivo*. Lower concentrations of *Shh* released from the notochord and the floorplate presumably induce motor neurons at more distant ventrolateral regions in a process that has been shown to be contact-independent *in vitro* (Yamada, T. *et al.* (1993) Cell 73:673-686). In explants taken at midbrain and forebrain levels, *Shh* also induces the appropriate ventrolateral neuronal cell types, dopaminergic (Heynes, M. *et al.* (1995) Neuron 15:35-44; Wang, M.Z. *et al.* (1995) Nature Med. 1:1184-1188) and cholinergic  
 30 (Ericson, J. *et al.* (1995) Cell 81:747-756) precursors, respectively, indicating that *Shh* is a common inducer of ventral specification over the entire length of the CNS. These

*Shh* from the midline also patterns the paraxial regions of the vertebrate embryo, the somites in the trunk (Fan *et al.* (1995) supra) and the head mesenchyme rostral of the somites (Hammerschmidt *et al.* (1996) supra). In chick and mouse paraxial mesoderm explants, *Shh* promotes the expression of sclerotome specific markers like *Pax1* and *Twist*, at the expense of the dermamyotomal marker *Pax3*. Moreover, filter barrier experiments suggest that *Shh* mediates the induction of the sclerotome directly rather than by activation of a secondary signaling mechanism (Fan, C.-M. and Tessier-Lavigne, M. (1994) Cell 79, 1175-1186).

*Shh* also induces myotomal gene expression (Hammerschmidt *et al.* (1996) supra; Johnson, R.L. *et al.* (1994) Cell 79:1165-1173; Münsterberg, A.E. *et al.* (1995) Genes Dev. 9:2911-2922; Weinberg, E.S. *et al.* (1996) Development 122:271-280), although recent experiments indicate that members of the WNT family, vertebrate homologues of *Drosophila wingless*, are required in concert (Münsterberg *et al.* (1995) supra). Puzzlingly, myotomal induction in chicks requires higher *Shh* concentrations than the induction of sclerotomal markers (Münsterberg *et al.* (1995) supra), although the sclerotome originates from somitic cells positioned much closer to the notochord. Similar results were obtained in the zebrafish, where high concentrations of Hedgehog induce myotomal and repress sclerotomal marker gene expression (Hammerschmidt *et al.* (1996) supra). In contrast to amniotes, however, these observations are consistent with the architecture of the fish embryo, as here, the myotome is the predominant and more axial component of the somites. Thus, modulation of *Shh* signaling and the acquisition of new signaling factors may have modified the somite structure during vertebrate evolution.

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*Patched* was originally identified in *Drosophila* as a segment polarity gene, one of a group of developmental genes that affect cell differentiation within the individual segments that occur in a homologous series along the anterior-posterior axis of the embryo. See Hooper, J.E. et al. (1989) Cell 59:751; and Nakano, Y. et al. (1989) Nature 341:508. Patterns of expression of the vertebrate homologue of *patched* suggest its involvement in the development of neural tube, skeleton, limbs, craniofacial structure, and skin.

Genetic and functional studies demonstrate that *patched* is part of the hedgehog signaling cascade, an evolutionarily conserved pathway that regulates expression of a number of downstream genes. See Perrimon, N. (1995) Cell 80:517; and Perrimon, N. (1996) Cell 86:513. *Patched* participates in the constitutive transcriptional repression of the target genes; its effect is opposed by a secreted glycoprotein, encoded by hedgehog, or a vertebrate homologue, which induces transcriptional activation. Genes under control of this pathway include members of the Wnt and TGF-beta families.

*Patched* proteins possess two large extracellular domains, twelve transmembrane segments, and several cytoplasmic segments. See Hooper, supra; Nakano, supra; Johnson, R.L. et al. (1996) Science 272:1668; and Hahn, H. et al. (1996) Cell 85:841. The biochemical role of *patched* in the hedgehog signaling pathway is unclear. Direct interaction with the hedgehog protein has, however, been reported (Chen, Y. et al. (1996) Cell 87:553), and *patched* may participate in a hedgehog receptor complex along with another transmembrane protein encoded by the *smoothed* gene. See Perrimon, supra; and Chen, supra.

The human homologue of *patched* was recently cloned and mapped to chromosome 9q22.3. See Johnson, supra; and Hahn, supra. This region has been implicated in basal cell nevus syndrome (BCNS), which is characterized by developmental abnormalities including rib and craniofacial alterations, abnormalities of the hands and feet, and spina bifida.

BCNS also predisposes to multiple tumor types, the most frequent being basal cell carcinomas (BCC) that occur in many locations on the body and appear within the first two decades of life. Most cases of BCC, however, are unrelated to the syndrome and arise sporadically in small numbers on sun-exposed sites of middle-aged or older people of northern European ancestry.

Recent studies in BCNS-related and sporadic BCC suggest that a functional loss of both alleles of *patched* leads to development of BCC. See Johnson, supra; Hahn, supra; and Gailani, M.R. et al. (1996) Nature Genetics 14:78. Single allele deletions of chromosome 9q22.3 occur frequently in both sporadic and hereditary BCC. Linkage analysis revealed that the defective inherited allele was retained and the normal allele was lost in tumors from BCNS patients.

Sporadic tumors also demonstrated a loss of both functional alleles of *patched*. Of twelve tumors in which *patched* mutations were identified with a single strand conformational polymorphism screening assay, nine had chromosomal deletion of the second allele and the other three had inactivating mutations in both alleles (Gailani, supra). The alterations did not occur in the corresponding germline DNA.

Most of the identified mutations resulted in premature stop codons or frame shifts. Lench, N.J., et al., *Hum. Genet.* 1997 Oct; 100(5-6): 497-502. Several, however, were point mutations leading to amino acid substitutions in either extracellular or cytoplasmic domains. These sites of mutation may indicate functional importance for interaction with extracellular proteins or with cytoplasmic members of the downstream signaling pathway.

The involvement of *patched* in the inhibition of gene expression and the occurrence of frequent allelic deletions of *patched* in BCC support a tumor suppressor function for this gene. Its role in the regulation of gene families known to be involved in cell signaling and intercellular communication provides a possible mechanism of tumor suppression.

### Summary of the Invention

One aspect of the present invention makes available methods and reagents for inhibiting *smoothened-dependent* pathway activation. In certain embodiments, the subject methods can be used to counteract the phenotypic effects of unwanted activation of a hedgehog pathway, such as resulting from *hedgehog* gain-of-function, *ptc* loss-of-function or *smoothened* gain-of-function mutations. For instance, the subject method can involve contacting a cell (*in vitro* or *in vivo*) with a *hedgehog* antagonist (defined infra), such as a

steroidal alkaloid or other small molecule in an amount sufficient to antagonize a smoothened-dependent pathway activation pathway.

Another aspect of the present invention makes available methods and reagents for activating *smoothened-dependent* pathway activation, e.g, to mimic all or certain of the effects that treatment with a hedgehog protein might cause. The subject method can involve contacting a cell (*in vitro* or *in vivo*) with a *smoothened* agonist (defined *infra*) in an amount sufficient to activate a smoothened-dependent pathway activation pathway.

The subject methods and compounds may be used to regulate proliferation and/or differentiation of cells *in vitro* and/or *in vivo*, e.g., in the formation of tissue from stem cells, or to prevent the growth of hyperproliferative cells to illustrate but a few uses.

The subject compounds may be formulated as a pharmaceutical preparation comprising a pharmaceutically acceptable excipient. Hedgehog antagonists of the invention and/or preparations comprising them may be administered to a patient to treat conditions involving unwanted cell proliferation, e.g., cancer and/or tumors (such as medullablastoma, basal cell carcinoma, etc.), non-malignant hyperproliferative disorders, etc. Smoothened agonists can also be used to regulate the growth and differentiation of normal tissues. In certain embodiments, such compounds or preparations are administered systemically and/or locally, e.g., topically.

## 20 Brief Description of the Drawings

**Figure 1.** Structures of the synthetic compounds AY 9944 and triparanol, of the plant steroidal alkaloids jervine, cyclopamine and tomatidine, and of cholesterol.

**Figure 2.** Holoprosencephaly induced in chick embryos exposed to jervine (4). (A) SEM of external facial features of an untreated embryo. (B, C, D and E) Embryos exposed to 10  $\mu$ M jervine with variable loss of midline tissue and resulting fusion of the paired, lateral olfactory processes (olf), optic vesicles (Opt), and maxillary (Mx) and mandibular (Mn) processes. A complete fusion of the optic vesicles (E) lead to true cyclopia.

**Figure 3.** Synthetic and plant derived teratogens block endogenous *Shh* signaling in explanted chick tissues (41). (A) Midline tissue was removed from stage 9-10 chick embryos at a level just rostral to Hensen's node (white dashed line), and further dissected

(black dashed lines) to yield an explant containing an endogenous source of *Shh* signal (notochord) and a responsive tissue (neural plate ectoderm). After two days of culture in a collagen gel matrix, the neural ectoderm expresses markers of floor plate cells (HNF3 $\beta$ , rhodamine) and motor neurons (Isl-1, FITC) in untreated control explants (B) and explants

5 cultured with the non-teratogenic alkaloid tomatidine (50  $\mu$ M, C). Intermedia doses of the teratogenic compounds AY 9944 (0.5  $\mu$ M, D), triparanol (0.25  $\mu$ M, E), jervine (0.5  $\mu$ M, F) and cyclopamine (0.25  $\mu$ M, G) block induction of HNF3 $\beta$ , which requires a high level of *Shh* pathway activation, while permitting induction of Isl-1, which requires a lower level of *Shh* pathway activation (see text). Higher doses of the teratogenic compounds AY

10 9944 (4.0  $\mu$ M, H), triparanol (1.0  $\mu$ M, I), jervine (4.0  $\mu$ M, J) and cyclopamine (1.0  $\mu$ M, K) and fully inhibit HNF3 $\beta$  and Isl-1 induction.

**Figure 4.** Teratogenic compounds do not inhibit *Shh* autoprocessing in vivo (47). Stably transfected HK293 cells expressing *Shh* protein under ecdysone-inducible control (lanes 1, 2, 3) were treated with jervine (lanes 4, 5) cyclopamine (lanes 6, 7), tomatidine

15 (lanes 10, 11), AY 9944 (lanes 12, 13) or triparanol (lanes 14, 15) and cell lysates were immunoblotted to assess the efficiency of autoprocessing. As seen in the untreated control (lane 3), *Shh* in treated cells is efficiently processed with little or no detectable accumulation of precursor protein (M<sub>r</sub>45 kD). The processed amino-terminal product (*Shh*-N<sub>p</sub>) is cell associated and migrates faster than *Shh*-N protein from the media of

20 cultured cells transfected with a construct carrying an open reading frame truncated after Gly<sub>198</sub> (lane 8; *Shh*-N<sub>p</sub> and *Shh*-N both loaded in lanes 9 and 17). This faster migration and the lack of detectable protein in the culture medium (not shown) indicate that *Shh*-N<sub>p</sub> from treated cells likely carries a sterol adduct. The slower migrating species resulting from tomatidine treatment is ~1.9 kD larger, suggestive of a minor inhibition of signal sequence

25 cleavage (see asterisk; lanes 10, 11). Immunoblotted actin for each lane is shown as a loading control.

**Figure 5.** Plant steroidal alkaloids do not inhibit or participate in *Hh* autoprocessing in vitro (5). (A) Coomassie blue-stained SDS-polyacrylamide gel showing in vitro autocleavage reactions of the bacterially expressed His<sub>6</sub>Hh-C protein (~29 kD)

30 incubated for 3 hours at 30° C with no sterol additions (lane 1) or 12  $\mu$ M cholesterol to stimulate the autoprocessing reaction and generate a ~25 kD Hh-C product (lanes 2-27 and a ~5 kD NH<sub>2</sub>-terminal product (not resolved on this gel). The addition of jervine (lanes 3-

6), cyclopamine (lanes 8-11) and tomatidine (lanes 13-16) does not interfere with autoproducting, even when added in 27-fold excess to cholesterol (lanes 6, 11 and 16). (B) Coomassie blue-stained SDS-polyacrylamide gel showing that the His<sub>6</sub>Hh-C autocleavage reaction does not proceed when carried out in the absence of sterol (lane 1), or in the presence of jervine (lanes 2-5), cyclopamine (lanes 6-9) and tomatidine (lanes 10-13), even at 324  $\mu$ M concentrations of these steroidal alkaloid (lanes 5, 9 and 13). (C) Coomassie blue-stained SDS polyacrylamide gel of His<sub>6</sub>Hh-C autocleavage reactions carried out in the absence of sterols (lane 1), with 50 mM dithiothreitol (lane 2), 12  $\mu$ M cholesterol (lane 3) 12  $\mu$ M 7 dehydrocholesterol (lane 4) 12  $\mu$ M desmosterol (lane 5), 12  $\mu$ M muristerone (lanes 9, 10). The 27-carbon cholesterol precursors (lanes 4-6) stimulate His<sub>6</sub>Hh-C autocleavage reactions carried out in the absence of sterols (lane 1), with 50 mM dithiothreitol (lane 2), 12  $\mu$ M cholesterol (lane 3) 12  $\mu$ M 7 dehydrocholesterol (lane 4) 12  $\mu$ M lathosterol (lane 6), 12 and 350  $\mu$ M lanosterol (lanes 7, 8) and 12 and 350  $\mu$ M muristerone (lanes 9, 10). The 27-carbon cholesterol precursors (lanes 4-6) stimulate His<sub>6</sub>Hh-C autoproducting as efficiently as cholesterol (lane 3). The amino-terminal product migrates as a ~7 kD species (lane 2) when generated in the presence of 50 mM dithiothreitol and as a ~5 kD species (lanes 3-6) with a sterol adduct. Lanosterol (lanes 7 and 8) and muristerone (lanes 9 and 10) do not stimulate autoproducting above background (lane 1).

**Figure 6.** Teratogenic compounds inhibit neural ectoderm response to exogenous Shh-N protein (41). (A) Intermediate neural plate ectoderm, free of notochord and other tissues, was dissected as shown (dashed lines) from stage 9-10 chick embryos at a level just rostral to Hensen's node (see figure 3A). (B) Explanted intermediate neural plate tissue cultured in a collagen gel matrix for 20 hours expresses the dorsal marker Pax7 (FITC) and not the floor plate marker HNF3 $\beta$  (Rhodamine). (C) Addition of recombinant, purified Shh-N at 2 nM suppresses Pax7 expression. (D) Markers of motor neuron (Isl-1, FITC) and floor plate cell (HNF3 $\beta$ , rhodamine) fates are induced upon explant culture for 40 hours in the presence of 6.25 nM Shh-N. (E) At 25 nM Shh-N, HNF3 $\beta$  expression expands at the expense of Isl-1 expression, which is lost. The repression of Pax7 expression by 2 nM Shh-N is inhibited by (F) 0.5  $\mu$ M AY 9944, (G) 0.25  $\mu$ M triparanol, (H) 0.125  $\mu$ M jervine and (I) 0.0625  $\mu$ M cyclopamine, but not by (J) 50  $\mu$ M tomatidine. Induction of HNF3 $\beta$  is blocked while induction of Isl-1 at 25 NM Shh-N is maintained or

expanded at intermediate levels of AY 9949 (1.0  $\mu$ M, K), triparanol (0.25  $\mu$ M, L), jervine (0.25  $\mu$ M, M), and cyclopamine (0.125  $\mu$ M, N). Tomatidine at 25 nM displays a slight inhibitory effect with decrease in HNF3 $\beta$  expression and an increase in the number Isl-1 expressing cells. HNF3 $\beta$  and Isl-1 induction are completely blocked at 2-fold higher doses of inhibitors AY 9944 (2.0  $\mu$ M, P), triparanol (0.5  $\mu$ M, Q), jervine (0.5,  $\mu$ M, R) and cyclopamine (0.25  $\mu$ M, S). Tomatidine at 50  $\mu$ M (T) markedly reduces HNF3 $\beta$  induction and enhances Isl-1 induction. Note that for each teratogenic compound the concentrations required to block complete the response to 2nM Shh-N (F-I) are lower than those required to block completely the response to 25 nM Shh-N (P-S). Also note that the response to 25 nM Shh-N is only partially inhibited (K-N) at concentrations of teratogen 2 fold lower than those required to block this response completely. See text for further comment.

**Figure 7.** Jervine does not inhibit neural ectoderm response to BMP7 (41). (A) Ventral neural plate ectoderm was dissected as shown (dashed lines) from stage 9-10 chick embryos at a level just rostral to Hensen's node (see Fig. 3A). (B) Ventral neural plate explants cultured for 24 hours in a collagen gel matrix do not give rise to any migratory cells that can be visualized by immunostaining for the HNK-1 antigen. (C) Addition of 100 ng/ml BMP7 induces formation of numerous HNK-1 positive cells that migrate out from the explant (borders outlined by white dashed line). (D) Induction of migratory HNK-1 positive cells by 100 ng/ml BMP7 is not inhibited by the presence of 10  $\mu$ M jervine, nor by addition of the other plant-derived compounds (10  $\mu$ M cyclopamine, 50  $\mu$ M tomatidine; data not shown).

**Figure 8.** Morphology and gene expression patterns of control and *Shh*<sup>-/-</sup> primary hair germs. (A,B) Normal-appearing hair genus consisting of an epithelial placode and adjacent mesenchymal condensate (arrows) were detected in skin of both control (A) and *Shh*<sup>-/-</sup> (B) embryos at 15.5 days of gestation (H&E staining). (C-H) Altered abundance of Shh target genes in hair germs in *Shh*<sup>-/-</sup> mouse skin. Expression of Glil (C,D), Ptcl. (E,F), and *BMP-4* (G,H) transcripts was examined in E 15.5 mouse skin using digoxigenin-labeled cRNA probes. Note virtual absence of Glil in both epithelial and mesenchymal components of the mutant hair germ and reduced mesenchymal Ptcl expression in *Shh*<sup>-/-</sup> skin.

**Figure 9.** Inhibition of hair follicle morphogenesis, but not biochemical differentiation, in *Shh*<sup>-/-</sup> mouse skin. (A,B) Advanced hair follicle development in skin from control (A) but not *Shh*<sup>-/-</sup> (B) embryos at 17.5 days of gestation (H&E staining). Note dermal papilla (arrow) surrounded by epithelial bulb of the largest hair follicle, and organizing mesenchymal aggregates (arrowheads) adjacent to invaginating tips of less mature follicles (A). In striking contrast, dermal papillae are not detected in *Shh* mutant skin (B). (C-F) Immunohistochemistry revealing similar patterns of keratin expression in control and *Shh*-deficient follicles. Absence of keratin K 14 immunostaining in subpopulation of keratinocytes in both control (C) and *Shh*<sup>-/-</sup> (D) hair follicles (arrows). Induction of non-epidermal keratin K17 in hair follicle keratinocytes in control (E) and *Shh* mutant (F) skin.

**Figure 10.** Impaired hair follicle development in *Shh* mutant skin grafted onto nude mice. (A) Gross appearance of nude mouse graft sites 6 weeks after transplantation. Note robust hair growth in control graft compared to hairless, but pigmented, *Shh*<sup>-/-</sup> skin graft. (B,C) H&E staining. Histologically normal-appearing skin in control graft (B) contains mature hair follicles with associated sebaceous glands and subcutaneous adipose tissue. Abnormal skin development in *Shh*<sup>-/-</sup> graft characterized by a thickened epidermis containing keratinocyte aggregates (arrows) at the base of the epidermis (C). (D-F) Immunohistochemistry. Unlike adjacent epidermal cells, *Shh*<sup>-/-</sup> keratinocyte aggregates do not express K5 (D, arrows) but are positive for Lef-1 localized to nuclei (E). Note also the presence of a small cluster of Lef-1 positive mesenchymal cells associated with the keratinocyte aggregate on the right (E). Immunostaining of abortive hair shafts with hair-specific keratin antibody AE 13 (F), revealing an advanced stage of follicle differentiation in *Shh* mutant skin.

**Figure 11.** Cyclopamine impairs vibrissa follicle morphogenesis in explant cultures. (A) vibrissa pad explants growing on Nuclepore membranes on day one and day eight in culture (dark-field). (B) FC-PCR analysis examining expression of transcripts encoding hair-specific markers *MHKAI* and *Hacl-1*, and an epidermal differentiation marker *filaggrin* (profilin). RNA was obtained from embryonic vibrissa pads when first isolated (Day 0) and after growth as explants (Day 7) in the presence or absence of 10<sup>-6</sup>M cyclopamine. Each lane contains reaction products for RNA isolated from an individual vibrissa pad. (C) Morphogenesis of vibrissa follicles is blocked by cyclopamine, an



inhibitor of Shh signaling. Cyclopamine was present in the medium for the duration of the experiment.

**Figure 12.** Ptc +/- MEFs incubated with *Shh*Np for 5 days.

**Figure 13.** Ptc -/- MEFs 23-1 cultured with cyclopamine for 3 days.

5 **Figure 14.** Ptc -/- MEFs 23-4 cultured with cyclopamine for 16 hours.

**Figure 15.** Ptc -/- MEFs 21-4 cultured with tomatidine for 16 hours.

### Detailed Description of the Invention

#### *I. Overview*

10 The present invention relates to the discovery that signal transduction pathways regulated by *hedgehog*, *patched* (*ptc*), *gli* and/or *smoothened* can be inhibited, at least in part, by small molecules. While not wishing to bound by any particular theory, the activation of a receptor may be the mechanism by which these agents act. For example, the ability of these agents to inhibit proliferation of *patched* loss-of-function (*ptc*<sup>lof</sup>) cells may  
15 be due to the ability of such molecules to interact with *hedgehog*, *patched*, or *smoothened*, or at least to interfere with the ability of those proteins to activate a *hedgehog*, *ptc*, and/or *smoothened*-mediated signal transduction pathway.

It is, therefore, specifically contemplated that these small molecules which interfere with aspects of *hedgehog*, *ptc*, or *smoothened* signal transduction activity will likewise be  
20 capable of inhibiting proliferation (or other biological consequences) in cells having a *patched* loss-of-function phenotype, a *hedgehog* gain-of-function phenotype, or a *smoothened* gain-of-function phenotype. In preferred embodiments, the subject inhibitors are organic molecules having a molecular weight less than 2500 amu, more preferably less than 1500 amu, and even more preferably less than 750 amu, and are capable of inhibiting  
25 at least some of the biological activities of hedgehog proteins, preferably specifically in target cells.

Thus, the methods of the present invention include the use of small molecules which agonize *ptc* inhibition of *hedgehog* signalling, such as by inhibiting activation of *smoothened* or downstream components of the signal pathway, in the regulation of repair  
30 and/or functional performance of a wide range of cells, tissues and organs having the

phenotype of *ptc* loss-of-function, *hedgehog* gain-of-function, or *smoothened* gain-of-function. For instance, the subject method has therapeutic and cosmetic applications ranging from regulation of neural tissues, bone and cartilage formation and repair, regulation of spermatogenesis, regulation of smooth muscle, regulation of lung, liver and other organs arising from the primitive gut, regulation of hematopoietic function, regulation of skin and hair growth, etc. Moreover, the subject methods can be performed on cells which are provided in culture (*in vitro*), or on cells in a whole animal (*in vivo*). See, for example, PCT publications WO 95/18856 and WO 96/17924 (the specifications of which are expressly incorporated by reference herein).

10 In a preferred embodiment, the subject method can be to treat epithelial cells having a phenotype of *ptc* loss-of-function, *hedgehog* gain-of-function, or *smoothened* gain-of-function. For instance, the subject method can be used in treating or preventing basal cell carcinoma or other *hedgehog* pathway-related disorders.

In another preferred embodiment, the subject method can be used as part of a treatment regimen for malignant medulloblastoma and other primary CNS malignant neuroectodermal tumors.

In another aspect, the present invention provides pharmaceutical preparations comprising, as an active ingredient, a *hedgehog* antagonist, *ptc* agonist, or *smoothened* antagonist such as described herein, formulated in an amount sufficient to inhibit, *in vivo*, proliferation or other biological consequences of *ptc* loss-of-function, *hedgehog* gain-of-function, or *smoothened* gain-of-function.

The subject treatments using *hedgehog* antagonists, *patched* agonists, or *smoothened* antagonists can be effective for both human and animal subjects. Animal subjects to which the invention is applicable extend to both domestic animals and livestock, raised either as pets or for commercial purposes. Examples are dogs, cats, cattle, horses, sheep, hogs, and goats.

## II. Definitions

For convenience, certain terms employed in the specification, examples, and appended claims are collected here.

The phrase "aberrant modification or mutation" of a gene refers to such genetic lesions as, for example, deletions, substitution or addition of nucleotides to a gene, as well as gross chromosomal rearrangements of the gene and/or abnormal methylation of the gene. Likewise, mis-expression of a gene refers to aberrant levels of transcription of the gene relative to those levels in a normal cell under similar conditions, as well as non-wild-type splicing of mRNA transcribed from the gene.

"Basal cell carcinomas" exist in a variety of clinical and histological forms such as nodular-ulcerative, superficial, pigmented, morphealike, fibroepithelioma and nevoid syndrome. Basal cell carcinomas are the most common cutaneous neoplasms found in humans. The majority of new cases of nonmelanoma skin cancers fall into this category.

"Burn wounds" refer to cases where large surface areas of skin have been removed or lost from an individual due to heat and/or chemical agents.

The term "carcinoma" refers to a malignant new growth made up of epithelial cells tending to infiltrate surrounding tissues and to give rise to metastases. Exemplary carcinomas include: "basal cell carcinoma", which is an epithelial tumor of the skin that, while seldom metastasizing, has potentialities for local invasion and destruction; "squamous cell carcinoma", which refers to carcinomas arising from squamous epithelium and having cuboid cells; "carcinosarcoma", which include malignant tumors composed of carcinomatous and sarcomatous tissues; "adenocystic carcinoma", carcinoma marked by cylinders or bands of hyaline or mucinous stroma separated or surrounded by nests or cords of small epithelial cells, occurring in the mammary and salivary glands, and mucous glands of the respiratory tract; "epidermoid carcinoma", which refers to cancerous cells which tend to differentiate in the same way as those of the epidermis; i.e., they tend to form prickle cells and undergo cornification; "nasopharyngeal carcinoma", which refers to a malignant tumor arising in the epithelial lining of the space behind the nose; and "renal cell carcinoma", which pertains to carcinoma of the renal parenchyma composed of tubular cells in varying arrangements. Other carcinomatous epithelial growths are "papillomas", which refers to benign tumors derived from epithelium and having a papillomavirus as a causative agent; and "epidermoidomas", which refers to a cerebral or meningeal tumor formed by inclusion of ectodermal elements at the time of closure of the neural groove.

The "corium" or "dermis" refers to the layer of the skin deep to the epidermis, consisting of a dense bed of vascular connective tissue, and containing the nerves and

terminal organs of sensation. The hair roots, and sebaceous and sweat glands are structures of the epidermis which are deeply embedded in the dermis.

"Dental tissue" refers to tissue in the mouth which is similar to epithelial tissue, for example gum tissue. The method of the present invention is useful for treating periodontal disease.

"Dermal skin ulcers" refer to lesions on the skin caused by superficial loss of tissue, usually with inflammation. Dermal skin ulcers which can be treated by the method of the present invention include decubitus ulcers, diabetic ulcers, venous stasis ulcers and arterial ulcers. Decubitus wounds refer to chronic ulcers that result from pressure applied to areas of the skin for extended periods of time. Wounds of this type are often called bedsores or pressure sores. Venous stasis ulcers result from the stagnation of blood or other fluids from defective veins. Arterial ulcers refer to necrotic skin in the area around arteries having poor blood flow.

The term "ED<sub>50</sub>" means the dose of a drug which produces 50% of its maximum response or effect.

An "effective amount" of, e.g., a *hedgehog* antagonist, with respect to the subject method of treatment, refers to an amount of the antagonist in a preparation which, when applied as part of a desired dosage regimen brings about, e.g., a change in the rate of cell proliferation and/or the state of differentiation of a cell and/or rate of survival of a cell according to clinically acceptable standards for the disorder to be treated or the cosmetic purpose.

The terms "epithelia", "epithelial" and "epithelium" refer to the cellular covering of internal and external body surfaces (cutaneous, mucous and serous), including the glands and other structures derived therefrom, e.g., corneal, esophageal, epidermal, and hair follicle epithelial cells. Other exemplary epithelial tissue includes: olfactory epithelium, which is the pseudostratified epithelium lining the olfactory region of the nasal cavity, and containing the receptors for the sense of smell; glandular epithelium, which refers to epithelium composed of secreting cells; squamous epithelium, which refers to epithelium composed of flattened plate-like cells. The term epithelium can also refer to transitional epithelium, like that which is characteristically found lining hollow organs that are subject

to great mechanical change due to contraction and distention, e.g., tissue which represents a transition between stratified squamous and columnar epithelium.

The term "epithelialization" refers to healing by the growth of epithelial tissue over a denuded surface.

- 5       The term "epidermal gland" refers to an aggregation of cells associated with the epidermis and specialized to secrete or excrete materials not related to their ordinary metabolic needs. For example, "sebaceous glands" are holocrine glands in the corium that secrete an oily substance and sebum. The term "sweat glands" refers to glands that secrete sweat, situated in the corium or subcutaneous tissue, opening by a duct on the body
- 10       surface.

- The term "epidermis" refers to the outermost and nonvascular layer of the skin, derived from the embryonic ectoderm, varying in thickness from 0.07-1.4 mm. On the palmar and plantar surfaces it comprises, from within outward, five layers: basal layer composed of columnar cells arranged perpendicularly; prickle-cell or spinous layer
- 15       composed of flattened polyhedral cells with short processes or spines; granular layer composed of flattened granular cells; clear layer composed of several layers of clear, transparent cells in which the nuclei are indistinct or absent; and horny layer composed of flattened, cornified non-nucleated cells. In the epidermis of the general body surface, the clear layer is usually absent.

- 20       "Excisional wounds" include tears, abrasions, cuts, punctures or lacerations in the epithelial layer of the skin and may extend into the dermal layer and even into subcutaneous fat and beyond. Excisional wounds can result from surgical procedures or from accidental penetration of the skin.

- The "growth state" of a cell refers to the rate of proliferation of the cell and/or the
- 25       state of differentiation of the cell. An "altered growth state" is a growth state characterized by an abnormal rate of proliferation, e.g., a cell exhibiting an increased or decreased rate of proliferation relative to a normal cell.

- The term "hair" refers to a threadlike structure, especially the specialized epidermal structure composed of keratin and developing from a papilla sunk in the corium, produced
- 30       only by mammals and characteristic of that group of animals. Also, "hair" may refer to the aggregate of such hairs. A "hair follicle" refers to one of the tubular-invaginations of the

epidermis enclosing the hairs, and from which the hairs grow. "Hair follicle epithelial cells" refers to epithelial cells which surround the dermal papilla in the hair follicle, e.g., stem cells, outer root sheath cells, matrix cells, and inner root sheath cells. Such cells may be normal non-malignant cells, or transformed/immortalized cells.

5       The term "*hedgehog* antagonist" refers to an agent which potentiates or recapitulates the bioactivity of *patched*, such as to repress transcription of target genes. Preferred *hedgehog* antagonists can be used to overcome a *ptc* loss-of-function and/or a *smoothened* gain-of-function, the latter also being referred to as *smoothened* antagonists. The term '*hedgehog* antagonist' as used herein refers not only to any agent that may act by  
10       directly inhibiting the normal function of the *hedgehog* protein, but also to any agent that inhibits the *hedgehog* signalling pathway, and thus recapitulates the function of *ptc*.

      The term "*hedgehog* gain-of-function" refers to an aberrant modification or mutation of a *ptc* gene, *hedgehog* gene, or *smoothened* gene, or a decrease (or loss) in the level of expression of such a gene, which results in a phenotype which resembles  
15       contacting a cell with a *hedgehog* protein, e.g., aberrant activation of a *hedgehog* pathway. The gain-of-function may include a loss of the ability of the *ptc* gene product to regulate the level of expression of Ci genes, e.g., *Gli1*, *Gli2*, and *Gli3*. The term '*hedgehog* gain-of-function' is also used herein to refer to any similar cellular phenotype (e.g., exhibiting excess proliferation) which occurs due to an alteration anywhere in the *hedgehog* signal  
20       transduction pathway, including, but not limited to, a modification or mutation of *hedgehog* itself. For example, a tumor cell with an abnormally high proliferation rate due to activation of the *hedgehog* signalling pathway would have a '*hedgehog* gain-of-function' phenotype, even if *hedgehog* is not mutated in that cell.

      As used herein, "immortalized cells" refers to cells which have been altered via  
25       chemical and/or recombinant means such that the cells have the ability to grow through an indefinite number of divisions in culture.

      "Internal epithelial tissue" refers to tissue inside the body which has characteristics similar to the epidermal layer in the skin. Examples include the lining of the intestine. The method of the present invention is useful for promoting the healing of certain internal  
30       wounds, for example wounds resulting from surgery.

The term "keratosis" refers to proliferative skin disorder characterized by hyperplasia of the horny layer of the epidermis. Exemplary keratotic disorders include keratosis follicularis, keratosis palmaris et plantaris, keratosis pharyngea, keratosis pilaris, and actinic keratosis.

5 The term "LD<sub>50</sub>" means the dose of a drug which is lethal in 50% of test subjects.

The term "nail" refers to the horny cutaneous plate on the dorsal surface of the distal end of a finger or toe.

The term "patched loss-of-function" refers to an aberrant modification or mutation of a *ptc* gene, or a decreased level of expression of the gene, which results in a phenotype which resembles contacting a cell with a hedgehog protein, e.g., aberrant activation of a hedgehog pathway. The loss-of-function may include a loss of the ability of the *ptc* gene product to regulate the level of expression of Ci genes, e.g., *Gli1*, *Gli2* and *Gli3*. The term 'ptc loss-of-function' is also used herein to refer to any similar cellular phenotype (e.g., exhibiting excess proliferation) which occurs due to an alteration anywhere in the hedgehog signal transduction pathway, including, but not limited to, a modification or mutation of *ptc* itself. For example, a tumor cell with an abnormally high proliferation rate due to activation of the hedgehog signalling pathway would have a 'ptc loss-of-function' phenotype, even if *ptc* is not mutated in that cell.

A "patient" or "subject" to be treated by the subject method can mean either a human or non-human animal.

The term "prodrug" is intended to encompass compounds which, under physiological conditions, are converted into the therapeutically active agents of the present invention. A common method for making a prodrug is to include selected moieties which are hydrolyzed under physiological conditions to reveal the desired molecule. In other embodiments, the prodrug is converted by an enzymatic activity of the host animal.

As used herein, "proliferating" and "proliferation" refer to cells undergoing mitosis.

Throughout this application, the term "proliferative skin disorder" refers to any disease/disorder of the skin marked by unwanted or aberrant proliferation of cutaneous tissue. These conditions are typically characterized by epidermal cell proliferation or incomplete cell differentiation, and include, for example, X-linked ichthyosis, psoriasis, atopic dermatitis, allergic contact dermatitis, epidermolytic hyperkeratosis, and seborrheic

dermatitis. For example, epidermodysplasia is a form of faulty development of the epidermis. Another example is "epidermolysis", which refers to a loosened state of the epidermis with formation of blebs and bullae either spontaneously or at the site of trauma.

As used herein, the term "psoriasis" refers to a hyperproliferative skin disorder which alters the skin's regulatory mechanisms. In particular, lesions are formed which involve primary and secondary alterations in epidermal proliferation, inflammatory responses of the skin, and an expression of regulatory molecules such as lymphokines and inflammatory factors. Psoriatic skin is morphologically characterized by an increased turnover of epidermal cells, thickened epidermis, abnormal keratinization, inflammatory cell infiltrates into the dermis layer and polymorphonuclear leukocyte infiltration into the epidermis layer resulting in an increase in the basal cell cycle. Additionally, hyperkeratotic and parakeratotic cells are present.

The term "skin" refers to the outer protective covering of the body, consisting of the corium and the epidermis, and is understood to include sweat and sebaceous glands, as well as hair follicle structures. Throughout the present application, the adjective "cutaneous" may be used, and should be understood to refer generally to attributes of the skin, as appropriate to the context in which they are used.

The term "*smoothened* gain-of-function" refers to an aberrant modification or mutation of a *smo* gene, or an increased level of expression of the gene, which results in a phenotype which resembles contacting a cell with a hedgehog protein, e.g., aberrant activation of a hedgehog pathway. While not wishing to be bound by any particular theory, it is noted that *ptc* may not signal directly into the cell, but rather interact with *smoothened*, another membrane-bound protein located downstream of *ptc* in *hedgehog* signaling (Marigo et al., (1996) Nature 384: 177-179). The gene *smo* is a segment-polarity gene required for the correct patterning of every segment in *Drosophila* (Alcedo et al., (1996) Cell 86: 221-232). Human homologs of *smo* have been identified. See, for example, Stone et al. (1996) Nature 384:129-134, and GenBank accession U84401. The *smoothened* gene encodes an integral membrane protein with characteristics of heterotrimeric G-protein-coupled receptors; i.e., 7-transmembrane regions. This protein shows homology to the *Drosophila* *Frizzled* (Fz) protein, a member of the *wingless* pathway. It was originally thought that *smo* encodes a receptor of the Hh signal. However, this suggestion was subsequently disproved, as evidence for *ptc* being the Hh receptor was obtained. Cells that



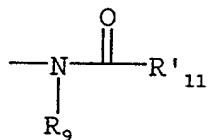
express *Smo* fail to bind Hh, indicating that *smo* does not interact directly with Hh (Nusse, (1996) Nature 384: 119-120). Rather, the binding of *Sonic hedgehog* (SHH) to its receptor, *PTCH*, is thought to prevent normal inhibition by *PTCH* of *smoothened* (SMO), a seven-span transmembrane protein.

- 5 Recently, it has been reported that activating *smoothened* mutations occur in sporadic basal cell carcinoma, Xie et al. (1998) Nature 391: 90-2, and primitive neuroectodermal tumors of the central nervous system, Reifenberger et al. (1998) Cancer Res 58: 1798-803.

- The term "therapeutic index" refers to the therapeutic index of a drug defined as  
10  $LD_{50}/ED_{50}$ .

- As used herein, "transformed cells" refers to cells which have spontaneously converted to a state of unrestrained growth, i.e., they have acquired the ability to grow through an indefinite number of divisions in culture. Transformed cells may be characterized by such terms as neoplastic, anaplastic and/or hyperplastic, with respect to  
15 their loss of growth control.

The term "acylamino" is art-recognized and refers to a moiety that can be represented by the general formula:



- 20 wherein  $R_9$  is as defined above, and  $R'_{11}$  represents a hydrogen, an alkyl, an alkenyl or  $-(CH_2)_m-R_8$ , where  $m$  and  $R_8$  are as defined above.

Herein, the term "aliphatic group" refers to a straight-chain, branched-chain, or cyclic aliphatic hydrocarbon group and includes saturated and unsaturated aliphatic groups, such as an alkyl group, an alkenyl group, and an alkynyl group.

- 25 The terms "alkenyl" and "alkynyl" refer to unsaturated aliphatic groups analogous in length and possible substitution to the alkyls described above, but that contain at least one double or triple bond respectively.

The terms "alkoxyl" or "alkoxy" as used herein refers to an alkyl group, as defined above, having an oxygen radical attached thereto. Representative alkoxyl groups include methoxy, ethoxy, propyloxy, tert-butoxy and the like. An "ether" is two hydrocarbons covalently linked by an oxygen. Accordingly, the substituent of an alkyl that renders that  
5 alkyl an ether is or resembles an alkoxyl, such as can be represented by one of -O-alkyl, -O-alkenyl, -O-alkynyl, -O-(CH<sub>2</sub>)<sub>m</sub>-R<sub>g</sub>, where m and R<sub>g</sub> are described above.

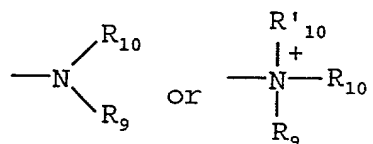
The term "alkyl" refers to the radical of saturated aliphatic groups, including straight-chain alkyl groups, branched-chain alkyl groups, cycloalkyl (alicyclic) groups, alkyl-substituted cycloalkyl groups, and cycloalkyl-substituted alkyl groups. In preferred  
10 embodiments, a straight chain or branched chain alkyl has 30 or fewer carbon atoms in its backbone (e.g., C<sub>1</sub>-C<sub>30</sub> for straight chains, C<sub>3</sub>-C<sub>30</sub> for branched chains), and more preferably 20 or fewer. Likewise, preferred cycloalkyls have from 3-10 carbon atoms in their ring structure, and more preferably have 5, 6 or 7 carbons in the ring structure.

Moreover, the term "alkyl" (or "lower alkyl") as used throughout the specification,  
15 examples, and claims is intended to include both "unsubstituted alkyls" and "substituted alkyls", the latter of which refers to alkyl moieties having substituents replacing a hydrogen on one or more carbons of the hydrocarbon backbone. Such substituents can include, for example, a halogen, a hydroxyl, a carbonyl (such as a carboxyl, an alkoxycarbonyl, a formyl, or an acyl), a thiocarbonyl (such as a thioester, a thioacetate, or a  
20 thioformate), an alkoxyl, a phosphoryl, a phosphate, a phosphonate, a phosphinate, an amino, an amido, an amidine, an imine, a cyano, a nitro, an azido, a sulfhydryl, an alkylthio, a sulfate, a sulfonate, a sulfamoyl, a sulfonamido, a sulfonyl, a heterocyclyl, an aralkyl, or an aromatic or heteroaromatic moiety. It will be understood by those skilled in the art that the moieties substituted on the hydrocarbon chain can themselves be  
25 substituted, if appropriate. For instance, the substituents of a substituted alkyl may include substituted and unsubstituted forms of amino, azido, imino, amido, phosphoryl (including phosphonate and phosphinate), sulfonyl (including sulfate, sulfonamido, sulfamoyl and sulfonate), and silyl groups, as well as ethers, alkylthios, carbonyls (including ketones, aldehydes, carboxylates, and esters), -CF<sub>3</sub>, -CN and the like. Exemplary substituted alkyls  
30 are described below. Cycloalkyls can be further substituted with alkyls, alkenyls, alkoxys, alkylthios, aminoalkyls, carbonyl-substituted alkyls, -CF<sub>3</sub>, -CN, and the like.

Unless the number of carbons is otherwise specified, "lower alkyl" as used herein means an alkyl group, as defined above, but having from one to ten carbons, more preferably from one to six carbon atoms in its backbone structure. Likewise, "lower alkenyl" and "lower alkynyl" have similar chain lengths. Throughout the application,  
 5 preferred alkyl groups are lower alkyls. In preferred embodiments, a substituent designated herein as alkyl is a lower alkyl.

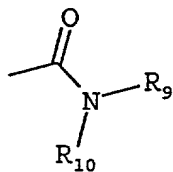
The term "alkylthio" refers to an alkyl group, as defined above, having a sulfur radical attached thereto. In preferred embodiments, the "alkylthio" moiety is represented by one of -S-alkyl, -S-alkenyl, -S-alkynyl, and -S-(CH<sub>2</sub>)<sub>m</sub>-R<sub>8</sub>, wherein m and R<sub>8</sub> are defined  
 10 above. Representative alkylthio groups include methylthio, ethylthio, and the like.

The terms "amine" and "amino" are art-recognized and refer to both unsubstituted and substituted amines, e.g., a moiety that can be represented by the general formula:



wherein R<sub>9</sub>, R<sub>10</sub> and R'<sub>10</sub> each independently represent a hydrogen, an alkyl, an alkenyl,  
 15 -(CH<sub>2</sub>)<sub>m</sub>-R<sub>8</sub>, or R<sub>9</sub> and R<sub>10</sub> taken together with the N atom to which they are attached complete a heterocycle having from 4 to 8 atoms in the ring structure; R<sub>8</sub> represents an aryl, a cycloalkyl, a cycloalkenyl, a heterocycle or a polycycle; and m is zero or an integer in the range of 1 to 8. In preferred embodiments, only one of R<sub>9</sub> or R<sub>10</sub> can be a carbonyl, e.g., R<sub>9</sub>, R<sub>10</sub> and the nitrogen together do not form an imide. In even more preferred  
 20 embodiments, R<sub>9</sub> and R<sub>10</sub> (and optionally R'<sub>10</sub>) each independently represent a hydrogen, an alkyl, an alkenyl, or -(CH<sub>2</sub>)<sub>m</sub>-R<sub>8</sub>. Thus, the term "alkylamine" as used herein means an amine group, as defined above, having a substituted or unsubstituted alkyl attached thereto, i.e., at least one of R<sub>9</sub> and R<sub>10</sub> is an alkyl group.

The term "amido" is art-recognized as an amino-substituted carbonyl and includes  
 25 a moiety that can be represented by the general formula:



wherein R<sub>9</sub>, R<sub>10</sub> are as defined above. Preferred embodiments of the amide will not include imides which may be unstable.

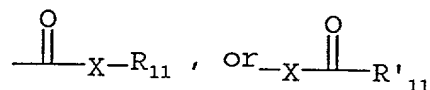
The term "aralkyl", as used herein, refers to an alkyl group substituted with an aryl group (e.g., an aromatic or heteroaromatic group).

The term "aryl" as used herein includes 5-, 6-, and 7-membered single-ring aromatic groups that may include from zero to four heteroatoms, for example, benzene, pyrrole, furan, thiophene, imidazole, oxazole, thiazole, triazole, pyrazole, pyridine, pyrazine, pyridazine and pyrimidine, and the like. Those aryl groups having heteroatoms in the ring structure may also be referred to as "aryl heterocycles" or "heteroaromatics." The aromatic ring can be substituted at one or more ring positions with such substituents as described above, for example, halogen, azide, alkyl, aralkyl, alkenyl, alkynyl, cycloalkyl, hydroxyl, alkoxyl, amino, nitro, sulfhydryl, imino, amido, phosphate, phosphonate, phosphinate, carbonyl, carboxyl, silyl, ether, alkylthio, sulfonyl, sulfonamido, ketone, aldehyde, ester, heterocyclyl, aromatic or heteroaromatic moieties, -CF<sub>3</sub>, -CN, or the like.

The term "aryl" also includes polycyclic ring systems having two or more cyclic rings in which two or more carbons are common to two adjoining rings (the rings are "fused rings") wherein at least one of the rings is aromatic, e.g., the other cyclic rings can be cycloalkyls, cycloalkenyls, cycloalkynyls, aryls and/or heterocyclyls.

The term "carbocycle", as used herein, refers to an aromatic or non-aromatic ring in which each atom of the ring is carbon.

The term "carbonyl" is art-recognized and includes such moieties as can be represented by the general formula:



wherein X is a bond or represents an oxygen or a sulfur, and R<sub>11</sub> represents a hydrogen, an alkyl, an alkenyl, -(CH<sub>2</sub>)<sub>m</sub>-R<sub>8</sub> or a pharmaceutically acceptable salt, R'<sub>11</sub> represents a

hydrogen, an alkyl, an alkenyl or  $-(CH_2)_m-R_8$ , where  $m$  and  $R_8$  are as defined above. Where  $X$  is an oxygen and  $R_{11}$  or  $R'_{11}$  is not hydrogen, the formula represents an "ester". Where  $X$  is an oxygen, and  $R_{11}$  is as defined above, the moiety is referred to herein as a carboxyl group, and particularly when  $R_{11}$  is a hydrogen, the formula represents a

5 "carboxylic acid". Where  $X$  is an oxygen, and  $R'_{11}$  is hydrogen, the formula represents a "formate". In general, where the oxygen atom of the above formula is replaced by sulfur, the formula represents a "thiocarbonyl" group. Where  $X$  is a sulfur and  $R_{11}$  or  $R'_{11}$  is not hydrogen, the formula represents a "thioester." Where  $X$  is a sulfur and  $R_{11}$  is hydrogen, the formula represents a "thiocarboxylic acid." Where  $X$  is a sulfur and  $R'_{11}$  is hydrogen,

10 the formula represents a "thiolformate." On the other hand, where  $X$  is a bond, and  $R_{11}$  is not hydrogen, the above formula represents a "ketone" group. Where  $X$  is a bond, and  $R_{11}$  is hydrogen, the above formula represents an "aldehyde" group.

The term "heteroatom" as used herein means an atom of any element other than carbon or hydrogen. Preferred heteroatoms are boron, nitrogen, oxygen, phosphorus, sulfur

15 and selenium.

The terms "heterocyclyl" or "heterocyclic group" refer to 3- to 10-membered ring structures, more preferably 3- to 7-membered rings, whose ring structures include one to four heteroatoms. Heterocycles can also be polycycles. Heterocyclyl groups include, for example, thiophene, thianthrene, furan, pyran, isobenzofuran, chromene, xanthene,

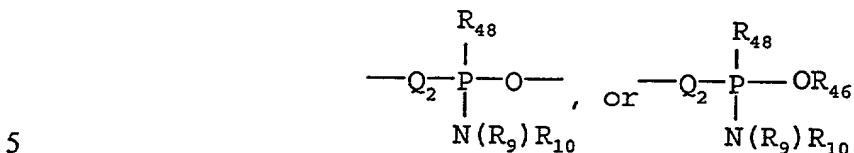
20 phenoxathiin, pyrrole, imidazole, pyrazole, isothiazole, isoxazole, pyridine, pyrazine, pyrimidine, pyridazine, indolizine, isoindole, indole, indazole, purine, quinolizine, isoquinoline, quinoline, phthalazine, naphthyridine, quinoxaline, quinazoline, cinnoline, pteridine, carbazole, carboline, phenanthridine, acridine, pyrimidine, phenanthroline, phenazine, phenarsazine, phenothiazine, furazan, phenoxazine, pyrrolidine, oxolane,

25 thiolane, oxazole, piperidine, piperazine, morpholine, lactones, lactams such as azetidinones and pyrrolidinones, sultams, sultones, and the like. The heterocyclic ring can be substituted at one or more positions with such substituents as described above, as for example, halogen, alkyl, aralkyl, alkenyl, alkynyl, cycloalkyl, hydroxyl, amino, nitro,

30 sulfhydryl, imino, amido, phosphate, phosphonate, phosphinate, carbonyl, carboxyl, silyl, ether, alkylthio, sulfonyl, ketone, aldehyde, ester, a heterocyclyl, an aromatic or heteroaromatic moiety,  $-CF_3$ ,  $-CN$ , or the like.

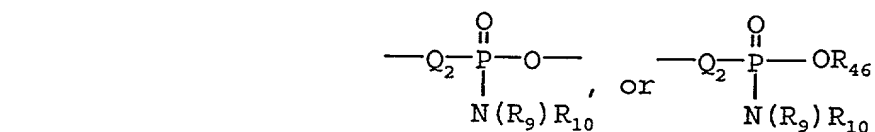
As used herein, the term "nitro" means -NO<sub>2</sub>; the term "halogen" designates -F, -Cl, -Br or -I; the term "sulfhydryl" means -SH; the term "hydroxyl" means -OH; and the term "sulfonyl" means -SO<sub>2</sub>-.

A "phosphonamidite" can be represented in the general formula:



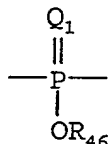
wherein R<sub>9</sub> and R<sub>10</sub> are as defined above, Q<sub>2</sub> represents O, S or N, and R<sub>48</sub> represents a lower alkyl or an aryl, Q<sub>2</sub> represents O, S or N.

A "phosphoramidite" can be represented in the general formula:



wherein R<sub>9</sub> and R<sub>10</sub> are as defined above, and Q<sub>2</sub> represents O, S or N.

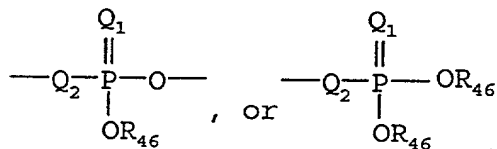
A "phosphoryl" can in general be represented by the formula:



wherein Q<sub>1</sub> represented S or O, and R<sub>46</sub> represents hydrogen, a lower alkyl or an aryl.

When used to substitute, for example, an alkyl, the phosphoryl group of the phosphorylalkyl can be represented by the general formula:

15



wherein Q<sub>1</sub> represented S or O, and each R<sub>46</sub> independently represents hydrogen, a lower alkyl or an aryl, Q<sub>2</sub> represents O, S or N. When Q<sub>1</sub> is an S, the phosphoryl moiety is a "phosphorothioate".

The terms "polycyclyl" or "polycyclic group" refer to two or more rings (e.g., cycloalkyls, cycloalkenyls, cycloalkynyls, aryls and/or heterocyclyls) in which two or more carbons are common to two adjoining rings, e.g., the rings are "fused rings". Rings that are joined through non-adjacent atoms are termed "bridged" rings. Each of the rings of the polycycle can be substituted with such substituents as described above, as for example, halogen, alkyl, aralkyl, alkenyl, alkynyl, cycloalkyl, hydroxyl, amino, nitro, sulfhydryl, imino, amido, phosphate, phosphonate, phosphinate, carbonyl, carboxyl, silyl, ether, alkylthio, sulfonyl, ketone, aldehyde, ester, a heterocyclyl, an aromatic or heteroaromatic moiety, -CF<sub>3</sub>, -CN, or the like.

The phrase "protecting group" as used herein means temporary substituents which protect a potentially reactive functional group from undesired chemical transformations. Examples of such protecting groups include esters of carboxylic acids, silyl ethers of alcohols, and acetals and ketals of aldehydes and ketones, respectively. The field of protecting group chemistry has been reviewed (Greene, T.W.; Wuts, P.G.M. *Protective Groups in Organic Synthesis*, 2<sup>nd</sup> ed.; Wiley: New York, 1991).

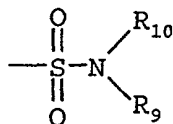
A "selenoalkyl" refers to an alkyl group having a substituted seleno group attached thereto. Exemplary "selenoethers" which may be substituted on the alkyl are selected from one of -Se-alkyl, -Se-alkenyl, -Se-alkynyl, and -Se-(CH<sub>2</sub>)<sub>m</sub>-R<sub>8</sub>, m and R<sub>8</sub> being defined above.

As used herein, the term "substituted" is contemplated to include all permissible substituents of organic compounds. In a broad aspect, the permissible substituents include acyclic and cyclic, branched and unbranched, carbocyclic and heterocyclic, aromatic and nonaromatic substituents of organic compounds. Illustrative substituents include, for example, those described herein above. The permissible substituents can be one or more and the same or different for appropriate organic compounds. For purposes of this invention, the heteroatoms such as nitrogen may have hydrogen substituents and/or any permissible substituents of organic compounds described herein which satisfy the valences of the heteroatoms. This invention is not intended to be limited in any manner by the permissible substituents of organic compounds.

It will be understood that "substitution" or "substituted with" includes the implicit proviso that such substitution is in accordance with permitted valence of the substituted

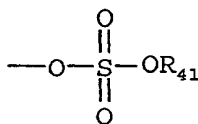
atom and the substituent, and that the substitution results in a stable compound, e.g., which does not spontaneously undergo transformation such as by rearrangement, cyclization, elimination, etc.

- The term "sulfamoyl" is art-recognized and includes a moiety that can be represented by the general formula:



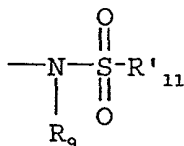
in which R<sub>9</sub> and R<sub>10</sub> are as defined above.

The term "sulfate" is art recognized and includes a moiety that can be represented by the general formula:



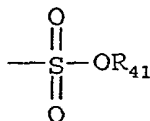
in which R<sub>41</sub> is as defined above.

The term "sulfonamido" is art recognized and includes a moiety that can be represented by the general formula:



- in which R<sub>9</sub> and R'<sub>11</sub> are as defined above.

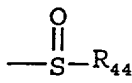
The term "sulfonate" is art-recognized and includes a moiety that can be represented by the general formula :



in which R<sub>41</sub> is an electron pair, hydrogen, alkyl, cycloalkyl, or aryl.



The terms "sulfoxido" or "sulfinyl", as used herein, refers to a moiety that can be represented by the general formula:



in which R<sub>44</sub> is selected from the group consisting of hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, heterocyclyl, aralkyl, or aryl.

Analogous substitutions can be made to alkenyl and alkynyl groups to produce, for example, aminoalkenyls, aminoalkynyls, amidoalkenyls, amidoalkynyls, iminoalkenyls, iminoalkynyls, thioalkenyls, thioalkynyls, carbonyl-substituted alkenyls or alkynyls.

As used herein, the definition of each expression, e.g., alkyl, m, n, etc., when it occurs more than once in any structure, is intended to be independent of its definition elsewhere in the same structure.

The terms triflyl, tosyl, mesyl, and nonafllyl are art-recognized and refer to trifluoromethanesulfonyl, *p*-toluenesulfonyl, methanesulfonyl, and nonafluorobutanesulfonyl groups, respectively. The terms triflate, tosylate, mesylate, and nonaflate are art-recognized and refer to trifluoromethanesulfonate ester, *p*-toluenesulfonate ester, methanesulfonate ester, and nonafluorobutanesulfonate ester functional groups and molecules that contain said groups, respectively.

The abbreviations Me, Et, Ph, Tf, Nf, Ts, Ms represent methyl, ethyl, phenyl, trifluoromethanesulfonyl, nonafluorobutanesulfonyl, *p*-toluenesulfonyl and methanesulfonyl, respectively. A more comprehensive list of the abbreviations utilized by organic chemists of ordinary skill in the art appears in the first issue of each volume of the *Journal of Organic Chemistry*; this list is typically presented in a table entitled Standard List of Abbreviations. The abbreviations contained in said list, and all abbreviations utilized by organic chemists of ordinary skill in the art are hereby incorporated by reference.

Certain compounds of the present invention may exist in particular geometric or stereoisomeric forms. The present invention contemplates all such compounds, including, cis- and trans-isomers, *R*- and *S*-enantiomers, diastereomers, (D)-isomers, (L)-isomers, the racemic mixtures thereof, and other mixtures thereof, as falling within the scope of the invention. Additional asymmetric carbon atoms may be present in a substituent such as an

alkyl group. All such isomers, as well as mixtures thereof, are intended to be included in this invention.

If, for instance, a particular enantiomer of a compound of the present invention is desired, it may be prepared by asymmetric synthesis, or by derivation with a chiral auxiliary, where the resulting diastereomeric mixture is separated and the auxiliary group cleaved to provide the pure desired enantiomers. Alternatively, where the molecule contains a basic functional group, such as amino, or an acidic functional group, such as carboxyl, diastereomeric salts may be formed with an appropriate optically active acid or base, followed by resolution of the diastereomers thus formed by fractional crystallization or chromatographic means well known in the art, and subsequent recovery of the pure enantiomers.

Contemplated equivalents of the compounds described above include compounds which otherwise correspond thereto, and which have the same general properties thereof (e.g., the ability to inhibit hedgehog signaling), wherein one or more simple variations of substituents are made which do not adversely affect the efficacy of the compound. In general, the compounds of the present invention may be prepared by the methods illustrated in the general reaction schemes as, for example, described below, or by modifications thereof, using readily available starting materials, reagents and conventional synthesis procedures. In these reactions, it is also possible to make use of variants which are in themselves known, but are not mentioned here.

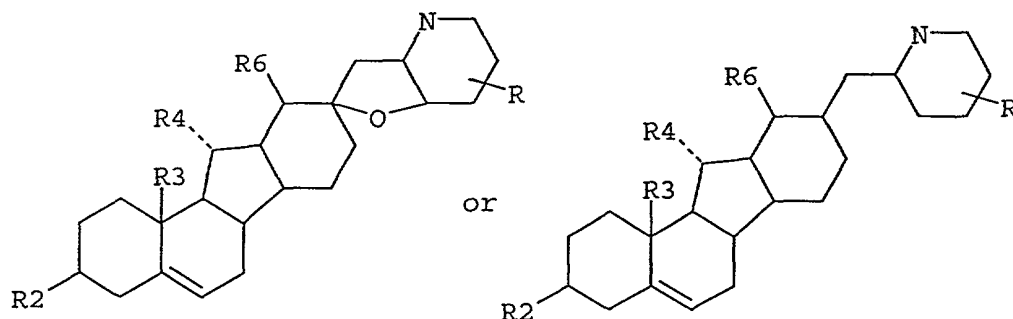
For purposes of this invention, the chemical elements are identified in accordance with the Periodic Table of the Elements, CAS version, Handbook of Chemistry and Physics, 67th Ed., 1986-87, inside cover. Also for purposes of this invention, the term "hydrocarbon" is contemplated to include all permissible compounds having at least one hydrogen and one carbon atom. In a broad aspect, the permissible hydrocarbons include acyclic and cyclic, branched and unbranched, carbocyclic and heterocyclic, aromatic and nonaromatic organic compounds which can be substituted or unsubstituted.

### III. Exemplary Compounds of the Invention.

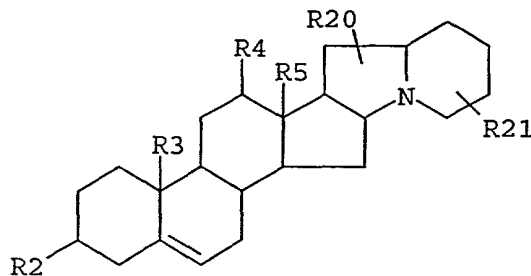
As described in further detail below, it is contemplated that the subject methods can be carried out using any of a variety of different steroidal alkaloids which can be

readily identified, e.g., by such drug screening assays as described herein. Steroidal alkaloids have a fairly complex nitrogen-containing nucleus. Two exemplary classes of steroidal alkaloids for use in the subject methods are the Solanum type and the Veratrum type. The above notwithstanding, in a preferred embodiment, the methods and compositions of the present invention make use of compounds having a steroidal alkaloid ring system of cyclopamine.

There are more than 50 naturally occurring veratrum alkaloids including veratramine, cyclopamine, cycloposine, jervine, and muldamine occurring in plants of the Veratrum spp. The Zigadenus spp., death camas, also produces several veratrum-type of steroidal alkaloids including zygacine. In general, many of the veratrum alkaloids (e.g., jervine, cyclopamine and cycloposine) consist of a modified steroid skeleton attached spiro to a furanopiperidine. A typical veratrum-type alkaloid may be represented by:

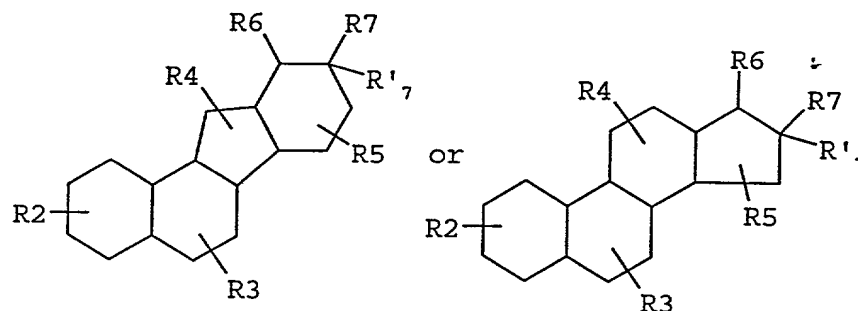


An example of the Solanum type is solanidine. This steroidal alkaloid is the nucleus (i.e., aglycone) for two important glycoalkaloids, solanine and chaconine, found in potatoes. Other plants in the Solanum family including various nightshades, Jerusalem cherries, and tomatoes also contain solanum-type glycoalkaloids. Glycoalkaloids are glycosides of alkaloids. A typical solanum-type alkaloid may be represented by:



Based on these structures, and the possibility that certain unwanted side effects can be reduced by some manipulation of the structure, a wide range of steroidal alkaloids are

contemplated as potential *hedgehog antagonists* for use in the subject method. For example, compounds useful in the subject methods include steroidal alkaloids represented in the general formulas (I), or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula I

wherein, as valence and stability permit,

$R_2$ ,  $R_3$ ,  $R_4$ , and  $R_5$ , represent one or more substitutions to the ring to which each is attached, for each occurrence, independently represent hydrogen, halogens, alkyls, alkenyls, alkynyls, aryls, hydroxyl, =O, =S, alkoxy, silyloxy, amino, nitro, thiol, amines, imines, amides, phosphoryls, phosphonates, phosphines, carbonyls, carboxyls, carboxamides, anhydrides, silyls, ethers, thioethers, alkylsulfonyls, arylsulfonyls, selenoethers, ketones, aldehydes, esters, sugar (e.g., monosaccharide, disaccharide, polysaccharide, etc.), carbamate (e.g., attached to the steroid at oxygen), carbonate, or -  
(CH<sub>2</sub>)<sub>m</sub>-R<sub>8</sub>;

$R_6$ ,  $R_7$ , and  $R'_7$ , are absent or represent, independently, halogens, alkyls, alkenyls, alkynyls, aryls, hydroxyl, =O, =S, alkoxy, silyloxy, amino, nitro, thiol, amines, imines, amides, phosphoryls, phosphonates, phosphines, carbonyls, carboxyls, carboxamides, anhydrides, silyls, ethers, thioethers, alkylsulfonyls, arylsulfonyls, selenoethers, ketones, aldehydes, esters, or -(CH<sub>2</sub>)<sub>m</sub>-R<sub>8</sub>, or

$R_6$  and  $R_7$ , or  $R_7$  and  $R'_7$ , taken together form a ring or polycyclic ring, e.g., which is substituted or unsubstituted,

with the proviso that at least one of  $R_6$ ,  $R_7$ , or  $R'_7$  is present and includes a primary or secondary amine;

R<sub>8</sub> represents an aryl, a cycloalkyl, a cycloalkenyl, a heterocycle, or a polycycle;  
and

m is an integer in the range 0 to 8 inclusive.

5 In preferred embodiments,

R<sub>2</sub> and R<sub>3</sub>, for each occurrence, is an -OH, alkyl, -O-alkyl, -C(O)-alkyl, or -C(O)-  
R<sub>8</sub>;

R<sub>4</sub>, for each occurrence, is an absent, or represents -OH, =O, alkyl, -O-alkyl, -  
C(O)-alkyl, or -C(O)-R<sub>8</sub>;

10 R<sub>6</sub>, R<sub>7</sub>, and R'<sub>7</sub> each independently represent, hydrogen, alkyls, alkenyls, alkynyls,  
amines, imines, amides, carbonyls, carboxyls, carboxamides, ethers, thioethers, esters, or -  
(CH<sub>2</sub>)<sub>m</sub>-R<sub>8</sub>, or

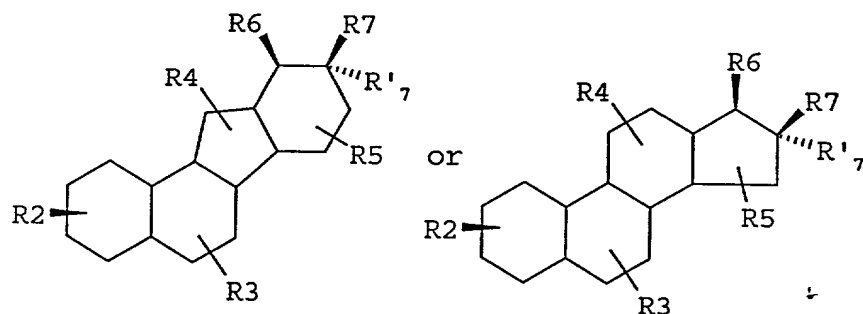
R<sub>7</sub>, and R'<sub>7</sub> taken together form a furanopiperidine, such as perhydrofuro[3,2-  
b]pyridine, a pyranopiperidine, a quinoline, an indole, a pyranopyrrole, a naphthyridine, a  
15 thiofuranopiperidine, or a thiopyranopiperidine

with the proviso that at least one of R<sub>6</sub>, R<sub>7</sub>, or R'<sub>7</sub> is present and includes a primary  
or secondary amine;

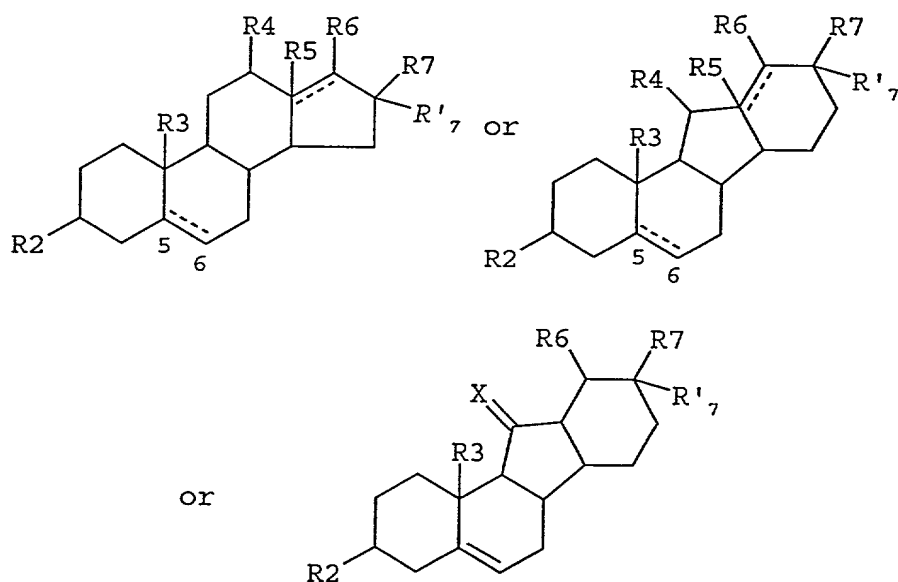
R<sub>8</sub> represents an aryl, a cycloalkyl, a cycloalkenyl, a heterocycle, or a polycycle,  
and preferably R<sub>8</sub> is a piperidine, pyrimidine, morpholine, thiomorpholine, pyridazine,

20

In certain preferred embodiments, the definitions outlined above apply, and the  
subject compounds are represented by general formula Ia or unsaturated forms thereof  
and/or seco-, nor- or homo-derivatives thereof:

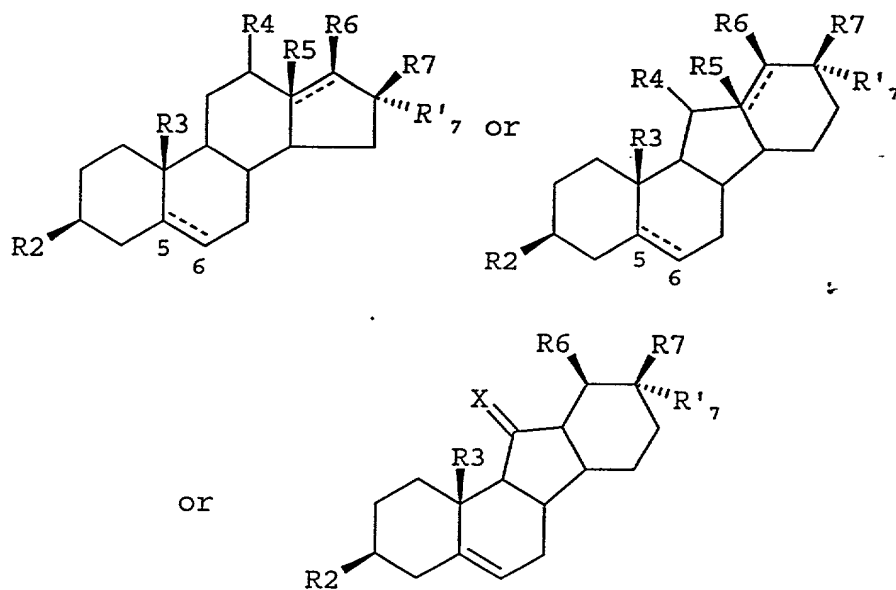
Formula Ia

In preferred embodiments, the subject hedgehog antagonists can be represented in one of the following general formulas (II) or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:

Formula II

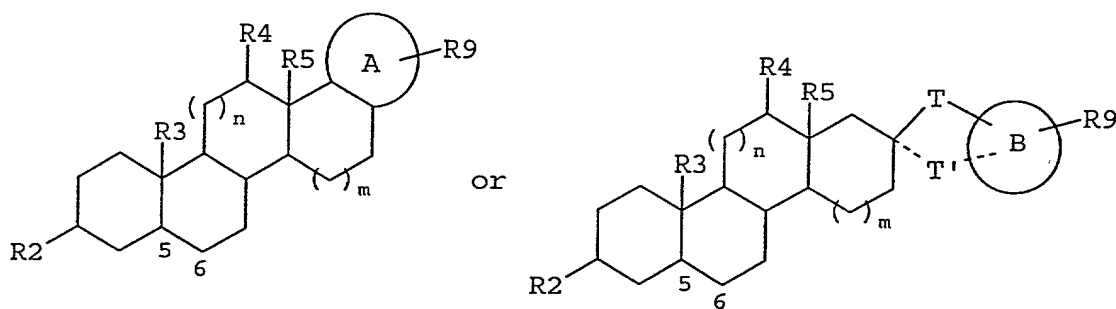
wherein  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ ,  $R_6$ ,  $R_7$ , and  $R'_7$  are as defined above, and X represents O or S, though preferably O.

In certain preferred embodiments, the definitions outlined above apply, and the subject compounds are represented by general formula IIa or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula IIa

In certain embodiments, the subject hedgehog antagonists are represented by the  
 5 general formula (III) or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula III

10 wherein

R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub> and R<sub>8</sub> are as defined above;

A and B represent monocyclic or polycyclic groups;

T represents an alkyl, an aminoalkyl, a carboxyl, an ester, an amide, ether or amine linkage of 1-10 bond lengths;

15 T' is absent, or represents an alkyl, an aminoalkyl, a carboxyl, an ester, an amide, ether or amine linkage of 1-3 bond lengths, wherein if T and T' are present together, than

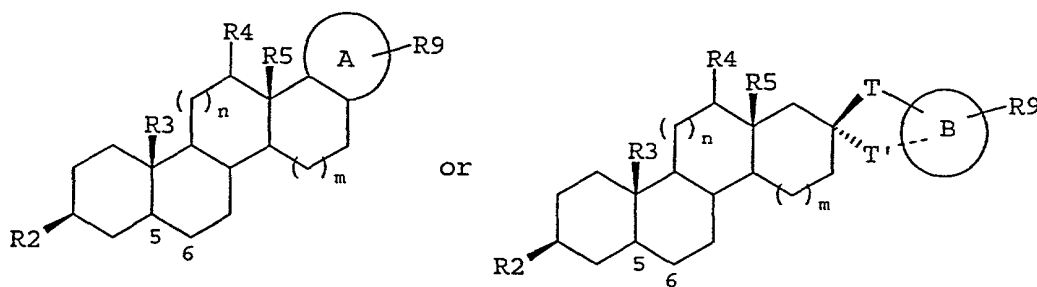
T and T' taken together with the ring A or B form a covalently closed ring of 5-8 ring atoms;

- R9 represents one or more substitutions to the ring A or B, which for each occurrence, independently represent halogens, alkyls, alkenyls, alkynyls, aryls, hydroxyl, =O, =S, alkoxyl, silyloxy, amino, nitro, thiol, amines, imines, amides, phosphoryls, phosphonates, phosphines, carbonyls, carboxyls, carboxamides, anhydrides, silyls, ethers, thioethers, alkylsulfonyls, arylsulfonyls, selenoethers, ketones, aldehydes, esters, or - (CH<sub>2</sub>)<sub>m</sub>-R<sub>8</sub>; and

n and m are, independently, zero, 1 or 2;

- with the proviso that A and R<sub>9</sub>, or T, T' B and R<sub>9</sub>, taken together include at least one primary or secondary amine.

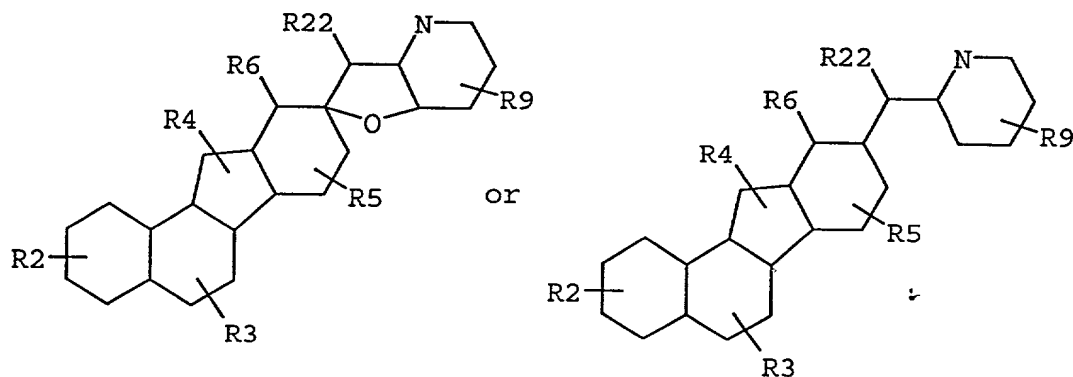
- In certain preferred embodiments, the definitions outlined above apply, and the subject compounds are represented by general formula IIIa or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula IIIa

- For example, the subject methods can utilize hedgehog antagonists based on the veratrum-type steroidal alkaloids jervine, cyclopamine, cycloposine, mukiamine or veratramine, e.g., which may be represented in the general formula (IV) or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:





Formula IV

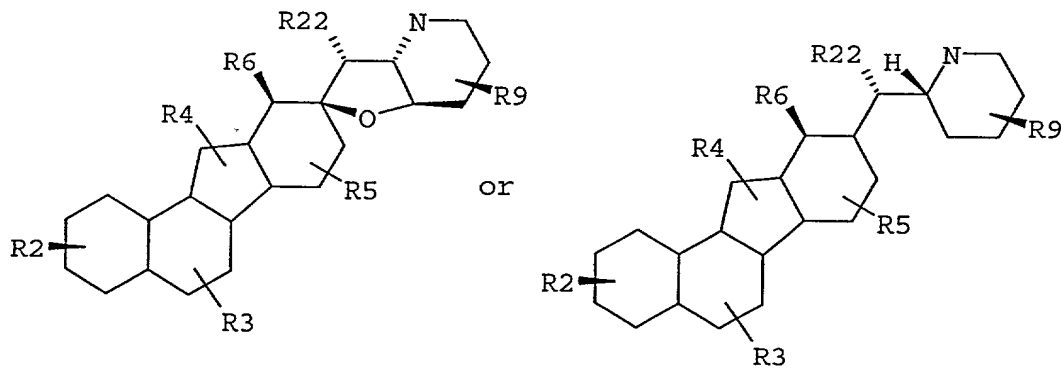
wherein

R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, R<sub>6</sub> and R<sub>9</sub> are as defined above;

5 R<sub>22</sub> is absent or represents an alkyl, an alkoxyl or -OH.

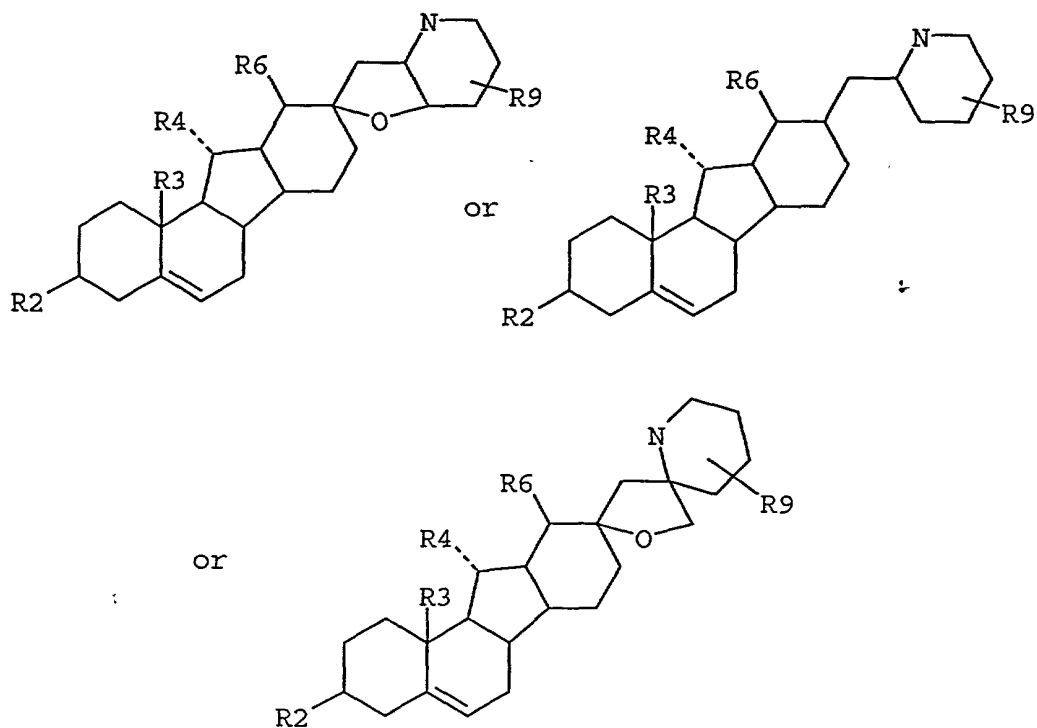
In certain preferred embodiments, the definitions outlined above apply, and the subject compounds are represented by general formula IVa or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:

10



Formula IVa

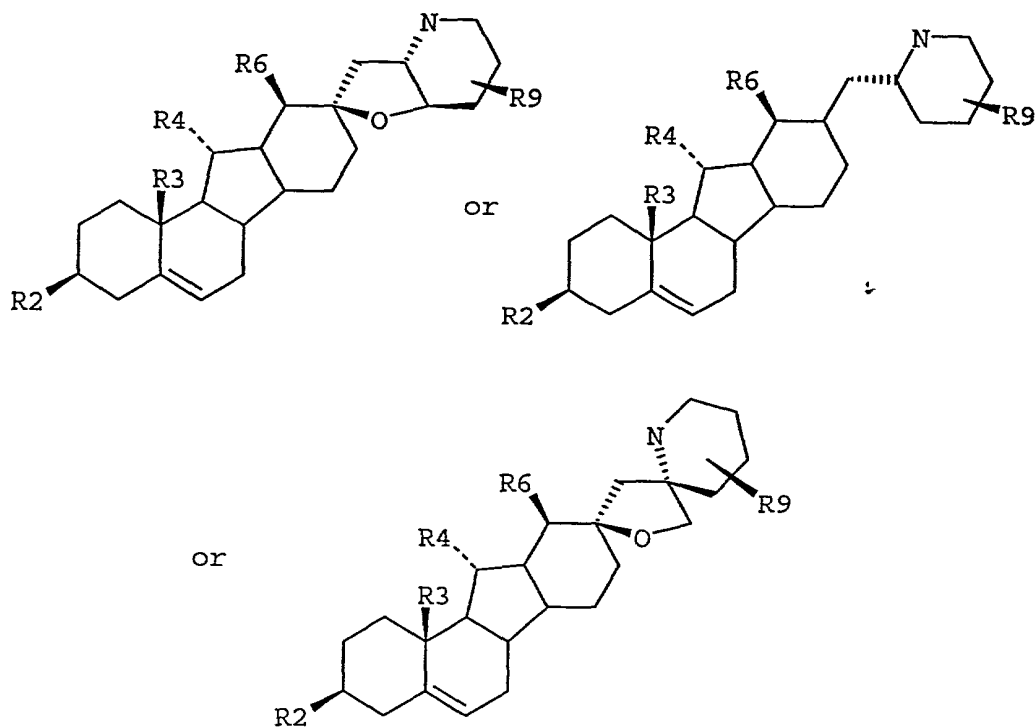
15 In even more preferred embodiments, the subject antagonists are represented in the formulas (V) or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula V

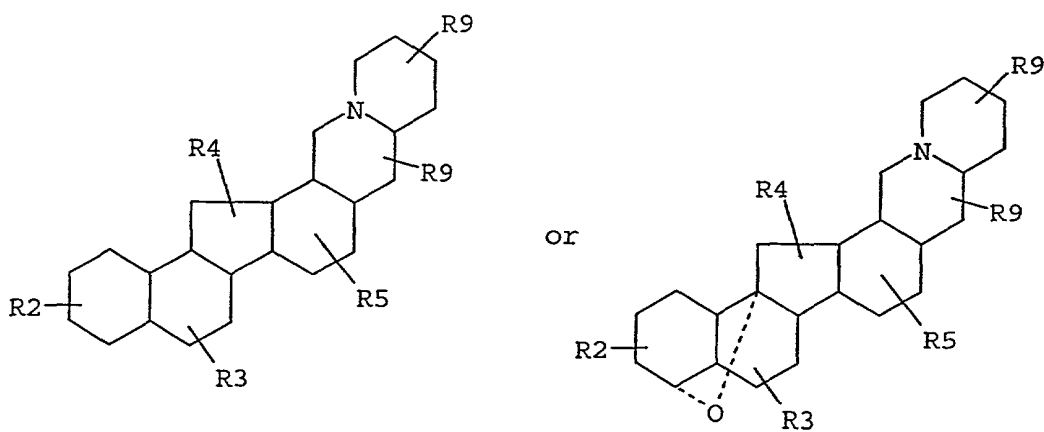
wherein R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>6</sub> and R<sub>9</sub> are as defined above;

- 5 In certain preferred embodiments, the definitions outlined above apply, and the subject compounds are represented by general formula Va or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula Va

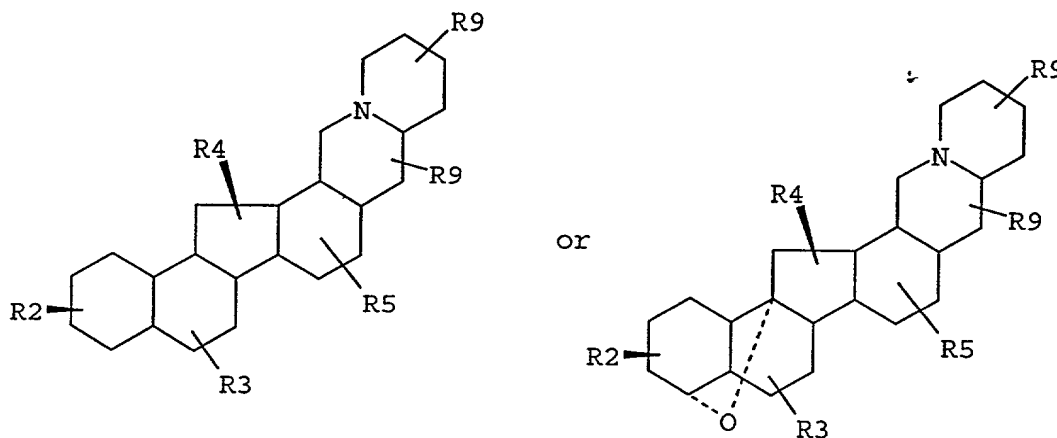
- Another class of hedgehog antagonists can be based on the veratrum-type steroidal  
 5 alkaloids resembling verticine and zygacine, e.g., represented in the general formulas (VI)  
 or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula VI

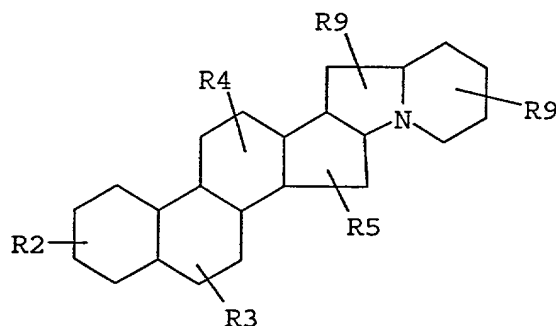
- 10 wherein R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub> and R<sub>9</sub> are as defined above.

In certain preferred embodiments, the definitions outlined above apply, and the subject compounds are represented by general formula VIa or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula VIa

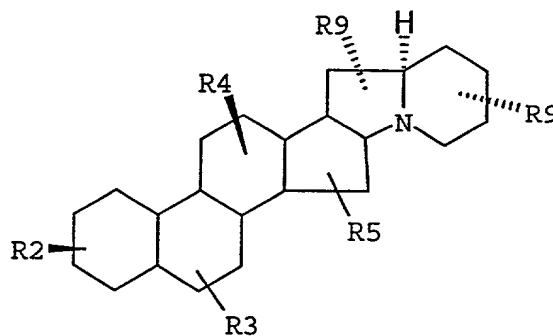
Still another class of potential hedgehog antagonists are based on the solanum-type steroidal alkaloids, e.g., solanidine, which may be represented in the general formula (VII) or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula VII

wherein R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub> and R<sub>9</sub> are as defined above.

15 In certain preferred embodiments, the definitions outlined above apply, and the subject compounds are represented by general formula VIIa or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula VIIa

In certain embodiments, the subject antagonists can be chosen on the basis of their selectivity for the hedgehog pathway. This selectivity can be for the hedgehog pathway versus other steroid-mediated pathways (such as testosterone or estrogen mediated activities), as well as selectivity for particular hedgehog pathways, e.g., which isotype specific for hedgehog (e.g., Shh, Ihh, Dhh) or the patched receptor (e.g., ptc-1, ptc-2). For instance, the subject method may employ steroidal alkaloids which do not substantially interfere with the biological activity of such steroids as aldosterone, androstane, androstene, androstenedione, androsterone, cholecalciferol, cholestane, cholic acid, corticosterone, cortisol, cortisol acetate, cortisone, cortisone acetate, deoxycorticosterone, digitoxigenin, ergocalciferol, ergosterol, estradiol-17- $\alpha$ , estradiol-17- $\beta$ , estriol, estrane, estrone, hydrocortisone, lanosterol, lithocholic acid, mestranol,  $\beta$ -methasone, prednisone, pregnane, pregnenolone, progesterone, spironolactone, testosterone, triamcinolone and their derivatives, at least so far as those activities are unrelated to ptc related signaling.

In one embodiment, the subject steroidal alkaloid for use in the present method has a  $K_d$  for members of the nuclear hormone receptor superfamily of greater than 1  $\mu$ M, and more preferably greater than 1 mM, e.g., it does not bind estrogen, testosterone receptors or the like. Preferably, the subject *hedgehog* antagonist has no estrogenic activity at physiological concentrations (e.g., in the range of 1 ng-1 mg/kg).

In this manner, untoward side effects which may be associated with certain members of the steroidal alkaloid class can be reduced. For example, using the drug screening assays described herein, the application of combinatorial and medicinal chemistry techniques to the steroidal alkaloids provides a means for reducing such unwanted negative side effects including personality changes, shortened life spans, cardiovascular diseases and vascular

occlusion, organ toxicity, hyperglycemia and diabetes, Cushnoid features, "wasting" syndrome, steroidal glaucoma, hypertension, peptic ulcers, and increased susceptibility to infections. For certain embodiments, it will be beneficial to reduce the teratogenic activity relative to jervine, as for example, in the use of the subject method to selectively inhibit spermatogenesis.

In a preferred embodiment, the subject antagonists are steroidal alkaloids other than spirosolane, tomatidine, jervine, etc.

In certain preferred embodiments, the subject inhibitors inhibit a hedgehog signal transduction pathway with an  $ED_{50}$  of 1mM or less, more preferably of 1  $\mu$ M or less, and even more preferably of 1nM or less.

In certain embodiments, the subject inhibitors inhibit a hedgehog signal transduction pathway with an  $ED_{50}$  of 1mM or less, more preferably 1  $\mu$ M or less, and even more preferably 1nM or less.

In particular embodiments, the steroidal alkaloid is chosen for use because it is more selective for one patched isoform over the next, e.g., 10-fold, and more preferably at least 100- or even 1000-fold more selective for one patched pathway (*ptc-1*, *ptc-2*) over another.

#### *IV. Exemplary Applications of Method and Compositions*

Another aspect of the present invention relates to a method of modulating a differentiated state, survival, and/or proliferation of a cell, such as a normal cell or a cell having a *ptc* loss-of-function, *hedgehog* gain-of-function, or *smoothened* gain-of-function, by contacting the cells with a compound as set forth above according to the subject method and as the circumstances may warrant.

For instance, it is contemplated by the invention that, in light of the findings of an apparently broad involvement of *hedgehog*, *ptc*, and *smoothened* in the formation of ordered spatial arrangements of differentiated tissues in vertebrates, the subject method could be used as part of a process for generating and/or maintaining an array of different, vertebrate tissue both *in vitro* and *in vivo*. The compound, whether inductive or anti-

inductive with respect proliferation or differentiation of a given tissue, can be, as appropriate, any of the preparations described above.

For example, the present method of using subject compounds is applicable to cell culture techniques wherein it is desirable to control the proliferation or differentiation of the cell. A subject compound may be employed in a method directed towards cells which have a *ptc* loss-of-function, *hedgehog* gain-of-function, or *smoothed* gain-of-function phenotype. *In vitro* neuronal culture systems have proved to be fundamental and indispensable tools for the study of neural development, as well as the identification of neurotrophic factors such as nerve growth factor (NGF), ciliary trophic factors (CNTF), and brain derived neurotrophic factor (BDNF). One use of the present method may be in cultures of neuronal stem cells, such as in the use of such cultures for the generation of new neurons and glia. In such embodiments of the subject method, the cultured cells can be contacted with a compound of the present invention in order to alter the rate of proliferation of neuronal stem cells in the culture and/or alter the rate of differentiation, or to maintain the integrity of a culture of certain terminally differentiated neuronal cells. In an exemplary embodiment, the subject method can be used to culture, for example, sensory neurons or, alternatively, motoneurons. Such neuronal cultures can be used as convenient assay systems as well as sources of implantable cells for therapeutic treatments.

According to the present invention, large numbers of non-tumorigenic neural progenitor cells can be perpetuated *in vitro* and their rate of proliferation and/or differentiation can be affected by contact with compounds of the present invention. Generally, a method is provided comprising the steps of isolating neural progenitor cells from an animal, perpetuating these cells *in vitro* or *in vivo*, preferably in the presence of growth factors, and regulating the differentiation of these cells into particular neural phenotypes, e.g., neurons and glia, by contacting the cells with a subject compound.

Progenitor cells are thought to be under a tonic inhibitory influence which maintains the progenitors in a suppressed state until their differentiation is required. However, recent techniques have been provided which permit these cells to be proliferated, and unlike neurons which are terminally differentiated and therefore non-dividing, they can be produced in unlimited number and are highly suitable for transplantation into heterologous and autologous hosts with neurodegenerative diseases.

By "progenitor" it is meant an oligopotent or multipotent stem cell which is able to divide without limit and, under specific conditions, can produce daughter cells which terminally differentiate such as into neurons and glia. These cells can be used for transplantation into a heterologous or autologous host. By heterologous is meant a host  
5 other than the animal from which the progenitor cells were originally derived. By autologous is meant the identical host from which the cells were originally derived.

Cells can be obtained from embryonic, post-natal, juvenile or adult neural tissue from any animal. By any animal is meant any multicellular animal which contains nervous tissue. More particularly, is meant any fish, reptile, bird, amphibian or mammal and the  
10 like. The most preferable donors are mammals, especially mice and humans.

In the case of a heterologous donor animal, the animal may be euthanized, and the brain and specific area of interest removed using a sterile procedure. Brain areas of particular interest include any area from which progenitor cells can be obtained which will serve to restore function to a degenerated area of the host's brain. These regions include  
15 areas of the central nervous system (CNS) including the cerebral cortex, cerebellum, midbrain, brainstem, spinal cord and ventricular tissue, and areas of the peripheral nervous system (PNS) including the carotid body and the adrenal medulla. More particularly, these areas include regions in the basal ganglia, preferably the striatum which consists of the caudate and putamen, or various cell groups such as the globus pallidus, the subthalamic  
20 nucleus, the nucleus basalis which is found to be degenerated in Alzheimer's Disease patients, or the substantia nigra pars compacta which is found to be degenerated in Parkinson's Disease patients.

Human heterologous neural progenitor cells may be derived from fetal tissue obtained from elective abortion, or from a post-natal, juvenile or adult organ donor.  
25 Autologous neural tissue can be obtained by biopsy, or from patients undergoing neurosurgery in which neural tissue is removed, in particular during epilepsy surgery, and more particularly during temporal lobectomies and hippocampalectomies.

Cells can be obtained from donor tissue by dissociation of individual cells from the connecting extracellular matrix of the tissue. Dissociation can be obtained using any  
30 known procedure, including treatment with enzymes such as trypsin, collagenase and the like, or by using physical methods of dissociation such as with a blunt instrument or by



mincing with a scalpel to allow outgrowth of specific cell types from a tissue. Dissociation of fetal cells can be carried out in tissue culture medium, while a preferable medium for dissociation of juvenile and adult cells is artificial cerebral spinal fluid (aCSF). Regular aCSF contains 124 mM NaCl, 5 mM KCl, 1.3 mM MgCl<sub>2</sub>, 2 mM CaCl<sub>2</sub>,  
 5 26 mM NaHCO<sub>3</sub>, and 10 mM D-glucose. Low Ca<sup>2+</sup> aCSF contains the same ingredients except for MgCl<sub>2</sub> at a concentration of 3.2 mM and CaCl<sub>2</sub> at a concentration of 0.1 mM.

Dissociated cells can be placed into any known culture medium capable of supporting cell growth, including MEM, DMEM, RPMI, F-12, and the like, containing supplements which are required for cellular metabolism such as glutamine and other  
 10 amino acids, vitamins, minerals and useful proteins such as transferrin and the like. Medium may also contain antibiotics to prevent contamination with yeast, bacteria and fungi such as penicillin, streptomycin, gentamicin and the like. In some cases, the medium may contain serum derived from bovine, equine, chicken and the like. A particularly preferable medium for cells is a mixture of DMEM and F-12.

15 Conditions for culturing should be close to physiological conditions. The pH of the culture media should be close to physiological pH, preferably between pH 6-8, more preferably close to pH 7, even more particularly about pH 7.4. Cells should be cultured at a temperature close to physiological temperature, preferably between 30 °C-40 °C, more preferably between 32 °C-38 °C, and most preferably between 35 °C-37 °C.

20 Cells can be grown in suspension or on a fixed substrate, but proliferation of the progenitors is preferably done in suspension to generate large numbers of cells by formation of "neurospheres" (see, for example, Reynolds et al. (1992) *Science* 255:1070-1709; and PCT Publications WO93/01275, WO94/09119, WO94/10292, and WO94/16718). In the case of propagating (or splitting) suspension cells, flasks are shaken  
 25 well and the neurospheres allowed to settle on the bottom corner of the flask. The spheres are then transferred to a 50 ml centrifuge tube and centrifuged at low speed. The medium is aspirated, the cells resuspended in a small amount of medium with growth factor, and the cells mechanically dissociated and resuspended in separate aliquots of media.

30 Cell suspensions in culture medium are supplemented with any growth factor which allows for the proliferation of progenitor cells and seeded in any receptacle capable

of sustaining cells, though as set out above, preferably in culture flasks or roller bottles. Cells typically proliferate within 3-4 days in a 37 °C incubator, and proliferation can be reinitiated at any time after that by dissociation of the cells and resuspension in fresh medium containing growth factors.

5 In the absence of substrate, cells lift off the floor of the flask and continue to proliferate in suspension forming a hollow sphere of undifferentiated cells. After approximately 3-10 days *in vitro*, the proliferating clusters (neurospheres) are fed every 2-7 days, and more particularly every 2-4 days by gentle centrifugation and resuspension in medium containing growth factor.

10 After 6-7 days *in vitro*, individual cells in the neurospheres can be separated by physical dissociation of the neurospheres with a blunt instrument, more particularly by triturating the neurospheres with a pipette. Single cells from the dissociated neurospheres are suspended in culture medium containing growth factors, and differentiation of the cells can be control in culture by plating (or resuspending) the cells in the presence of a subject  
15 compound.

To further illustrate other uses of the subject compounds, it is noted that intracerebral grafting has emerged as an additional approach to central nervous system therapies. For example, one approach to repairing damaged brain tissues involves the transplantation of cells from fetal or neonatal animals into the adult brain (Dunnett et al.  
20 (1987) *J Exp Biol* 123:265-289; and Freund et al. (1985) *J Neurosci* 5:603-616). Fetal neurons from a variety of brain regions can be successfully incorporated into the adult brain, and such grafts can alleviate behavioral defects. For example, movement disorder induced by lesions of dopaminergic projections to the basal ganglia can be prevented by grafts of embryonic dopaminergic neurons. Complex cognitive functions that are impaired  
25 after lesions of the neocortex can also be partially restored by grafts of embryonic cortical cells. The subject method can be used to regulate the growth state in the culture, or where fetal tissue is used, especially neuronal stem cells, can be used to regulate the rate of differentiation of the stem cells.

Stem cells useful in the present invention are generally known. For example,  
30 several neural crest cells have been identified, some of which are multipotent and likely represent uncommitted neural crest cells, and others of which can generate only one type

of cell, such as sensory neurons, and likely represent committed progenitor cells. The role of compounds employed in the present method to culture such stem cells can be to regulate differentiation of the uncommitted progenitor, or to regulate further restriction of the developmental fate of a committed progenitor cell towards becoming a terminally differentiated neuronal cell. For example, the present method can be used *in vitro* to regulate the differentiation of neural crest cells into glial cells, schwann cells, chromaffin cells, cholinergic sympathetic or parasympathetic neurons, as well as peptidergic and serotonergic neurons. The subject compounds can be used alone, or can be used in combination with other neurotrophic factors which act to more particularly enhance a particular differentiation fate of the neuronal progenitor cell.

In addition to the implantation of cells cultured in the presence of the subject compounds, yet another aspect of the present invention concerns the therapeutic application of a subject compound to regulate the growth state of neurons and other neuronal cells in both the central nervous system and the peripheral nervous system. The ability of *ptc*, *hedgehog*, and *smoothed* to regulate neuronal differentiation during development of the nervous system and also presumably in the adult state indicates that, in certain instances, the subject compounds can be expected to facilitate control of adult neurons with regard to maintenance, functional performance, and aging of normal cells; repair and regeneration processes in chemically or mechanically lesioned cells; and treatment of degeneration in certain pathological conditions. In light of this understanding, the present invention specifically contemplates applications of the subject method to the treatment protocol of (prevention and/or reduction of the severity of) neurological conditions deriving from: (i) acute, subacute, or chronic injury to the nervous system, including traumatic injury, chemical injury, vascular injury and deficits (such as the ischemia resulting from stroke), together with infectious/inflammatory and tumor-induced injury; (ii) aging of the nervous system including Alzheimer's disease; (iii) chronic neurodegenerative diseases of the nervous system, including Parkinson's disease, Huntington's chorea, amyotrophic lateral sclerosis and the like, as well as spinocerebellar degenerations; and (iv) chronic immunological diseases of the nervous system or affecting the nervous system, including multiple sclerosis.

As appropriate, the subject method can also be used in generating nerve prostheses for the repair of central and peripheral nerve damage. In particular, where a crushed or

severed axon is intubulated by use of a prosthetic device, subject compounds can be added to the prosthetic device to regulate the rate of growth and regeneration of the dendritic processes. Exemplary nerve guidance channels are described in U.S. patents 5,092,871 and 4,955,892.

- 5 In another embodiment, the subject method can be used in the treatment of neoplastic or hyperplastic transformations such as may occur in the central nervous system. For instance, the subject compounds can be utilized to cause such transformed cells to become either post-mitotic or apoptotic. The present method may, therefore, be used as part of a treatment for, e.g., malignant gliomas, meningiomas, medulloblastomas, neuroectodermal tumors, and ependymomas.

In a preferred embodiment, the subject method can be used as part of a treatment regimen for malignant medulloblastoma and other primary CNS malignant neuroectodermal tumors.

- 15 In certain embodiments, the subject method is used as part of treatment program for medulloblastoma. Medulloblastoma, a primary brain tumor, is the most common brain tumor in children. A medulloblastoma is a primitive neuroectodermal tumor arising in the posterior fossa. They account for approximately 25% of all pediatric brain tumors (Miller). Histologically, they are small round cell tumors commonly arranged in true rosettes, but may display some differentiation to astrocytes, ependymal cells or neurons (Rorke; Kleihues). PNET's may arise in other areas of the brain including the pineal gland (pineoblastoma) and cerebrum. Those arising in the supratentorial region generally fare worse than their PF counterparts.

- 25 Medulloblastoma/PNET's are known to recur anywhere in the CNS after resection, and can even metastasize to bone. Pretreatment evaluation should therefore include an examination of the spinal cord to exclude the possibility of "dropped metastases". Gadolinium-enhanced MRI has largely replaced myelography for this purpose, and CSF cytology is obtained postoperatively as a routine procedure.

- 30 In other embodiments, the subject method is used as part of treatment program for ependymomas. Ependymomas account for approximately 10% of the pediatric brain tumors in children. Grossly, they are tumors that arise from the ependymal lining of the ventricles and microscopically form rosettes, canals, and perivascular rosettes. In the

CHOP series of 51 children reported with ependymomas,  $\frac{3}{4}$  were histologically benign. Approximately  $\frac{2}{3}$  arose from the region of the 4th ventricle. One third presented in the supratentorial region. Age at presentation peaks between birth and 4 years, as demonstrated by SEER data as well as data from CHOP. The median age is about 5 years.

- 5 Because so many children with this disease are babies, they often require multimodal therapy.

Yet another aspect of the present invention concerns the observation in the art that *ptc*, *hedgehog*, and/or *smoothened* are involved in morphogenic signals involved in other vertebrate organogenic pathways in addition to neuronal differentiation as described above, having apparent roles in other endodermal patterning, as well as both mesodermal and endodermal differentiation processes. Thus, it is contemplated by the invention that compositions comprising one or more of the subject compounds can also be utilized for both cell culture and therapeutic methods involving generation and maintenance of non-neuronal tissue.

- 15 In one embodiment, the present invention makes use of the discovery that *ptc*, *hedgehog*, and *smoothened* are apparently involved in controlling the development of stem cells responsible for formation of the digestive tract, liver, lungs, and other organs which derive from the primitive gut. *Shh* serves as an inductive signal from the endoderm to the mesoderm, which is critical to gut morphogenesis. Therefore, for example, compounds of the instant method can be employed for regulating the development and maintenance of an artificial liver which can have multiple metabolic functions of a normal liver. In an exemplary embodiment, the subject method can be used to regulate the proliferation and differentiation of digestive tube stem cells to form hepatocyte cultures which can be used to populate extracellular matrices, or which can be encapsulated in biocompatible polymers, to form both implantable and extracorporeal artificial livers.

In another embodiment, therapeutic compositions of subject compounds can be utilized in conjunction with transplantation of such artificial livers, as well as embryonic liver structures, to regulate uptake of intraperitoneal implantation, vascularization, and *in vivo* differentiation and maintenance of the engrafted liver tissue.

- 30 In yet another embodiment, the subject method can be employed therapeutically to regulate such organs after physical, chemical or pathological insult. For instance,

therapeutic compositions comprising subject compounds can be utilized in liver repair subsequent to a partial hepatectomy.

The generation of the pancreas and small intestine from the embryonic gut depends on intercellular signalling between the endodermal and mesodermal cells of the gut. In particular, the differentiation of intestinal mesoderm into smooth muscle has been suggested to depend on signals from adjacent endodermal cells. One candidate mediator of endodermally derived signals in the embryonic hindgut is *Sonic hedgehog*. See, for example, Apelqvist et al. (1997) Curr Biol 7:801-4. The *Shh* gene is expressed throughout the embryonic gut endoderm with the exception of the pancreatic bud endoderm, which instead expresses high levels of the homeodomain protein *Ipfl/Pdx1* (insulin promoter factor 1/pancreatic and duodenal homeobox 1), an essential regulator of early pancreatic development. Apelqvist et al., supra, have examined whether the differential expression of *Shh* in the embryonic gut tube controls the differentiation of the surrounding mesoderm into specialised mesoderm derivatives of the small intestine and pancreas. To test this, they used the promoter of the *Ipfl/Pdx1* gene to selectively express *Shh* in the developing pancreatic epithelium. In *Ipfl/Pdx1-Shh* transgenic mice, the pancreatic mesoderm developed into smooth muscle and interstitial cells of Cajal, characteristic of the intestine, rather than into pancreatic mesenchyme and spleen. Also, pancreatic explants exposed to *Shh* underwent a similar program of intestinal differentiation. These results provide evidence that the differential expression of endodermally derived *Shh* controls the fate of adjacent mesoderm at different regions of the gut tube.

In the context of the present invention, it is contemplated therefore that the subject compounds can be used to control or regulate the proliferation and/or differentiation of pancreatic tissue both *in vivo* and *in vitro*.

There are a wide variety of pathological cell proliferative and differentiative conditions for which the inhibitors of the present invention may provide therapeutic benefits, with the general strategy being, for example, the correction of aberrant insulin expression, or modulation of differentiation. More generally, however, the present invention relates to a method of inducing and/or maintaining a differentiated state, enhancing survival and/or affecting proliferation of pancreatic cells, by contacting the cells with the subject inhibitors. For instance, it is contemplated by the invention that, in light of the apparent involvement of *ptc*, *hedgehog*, and *smoothed* in the formation of ordered

spatial arrangements of pancreatic tissues, the subject method could be used as part of a technique to generate and/or maintain such tissue both *in vitro* and *in vivo*. For instance, modulation of the function of *hedgehog* can be employed in both cell culture and therapeutic methods involving generation and maintenance  $\beta$ -cells and possibly also for  
5 non-pancreatic tissue, such as in controlling the development and maintenance of tissue from the digestive tract, spleen, lungs, urogenital organs (e.g., bladder), and other organs which derive from the primitive gut.

In an exemplary embodiment, the present method can be used in the treatment of hyperplastic and neoplastic disorders effecting pancreatic tissue, particularly those  
10 characterized by aberrant proliferation of pancreatic cells. For instance, pancreatic cancers are marked by abnormal proliferation of pancreatic cells which can result in alterations of insulin secretory capacity of the pancreas. For instance, certain pancreatic hyperplasias, such as pancreatic carcinomas, can result in hypoinsulinemia due to dysfunction of  $\beta$ -cells or decreased islet cell mass. To the extent that aberrant *ptc*, *hedgehog*, and *smoothened*  
15 signaling may be indicated in disease progression, the subject regulators can be used to enhance regeneration of the tissue after anti-tumor therapy.

Moreover, manipulation of *hedgehog* signaling properties at different points may be useful as part of a strategy for reshaping/repairing pancreatic tissue both *in vivo* and *in vitro*. In one embodiment, the present invention makes use of the apparent involvement of  
20 *ptc*, *hedgehog*, and *smoothened* in regulating the development of pancreatic tissue. In general, the subject method can be employed therapeutically to regulate the pancreas after physical, chemical or pathological insult. In yet another embodiment, the subject method can be applied to cell culture techniques, and in particular, may be employed to enhance the initial generation of prosthetic pancreatic tissue devices. Manipulation of proliferation  
25 and differentiation of pancreatic tissue, for example, by altering *hedgehog* activity, can provide a means for more carefully controlling the characteristics of a cultured tissue. In an exemplary embodiment, the subject method can be used to augment production of prosthetic devices which require  $\beta$ -islet cells, such as may be used in the encapsulation devices described in, for example, the Aebischer et al. U.S. Patent No. 4,892,538, the  
30 Aebischer et al. U.S. Patent No. 5,106,627, the Lim U.S. Patent No. 4,391,909, and the Sefton U.S. Patent No. 4,353,888. Early progenitor cells to the pancreatic islets are multipotential, and apparently coactivate all the islet-specific genes from the time they first

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cancer, etc.), *Shh*-induced tumors (e.g., lung cancer, chondrosarcomas, etc.), and other tumors (e.g., breast cancer, urogenital cancer (e.g., kidney, bladder, ureter, prostate, etc.), adrenal cancer, gastrointestinal cancer (e.g., stomach, intestine, etc.), etc.).

5 In still another embodiment of the present invention, compositions comprising one or more of the subject compounds can be used in the *in vitro* generation of skeletal tissue, such as from skeletogenic stem cells, as well as the *in vivo* treatment of skeletal tissue deficiencies. The present invention particularly contemplates the use of subject compounds to regulate the rate of chondrogenesis and/or osteogenesis. By "skeletal tissue deficiency", it is meant a deficiency in bone or other skeletal connective tissue at any site where it is  
10 desired to restore the bone or connective tissue, no matter how the deficiency originated, e.g. whether as a result of surgical intervention, removal of tumor, ulceration, implant, fracture, or other traumatic or degenerative conditions.

For instance, the method of the present invention can be used as part of a regimen for restoring cartilage function to a connective tissue. Such methods are useful in, for  
15 example, the repair of defects or lesions in cartilage tissue which is the result of degenerative wear such as that which results in arthritis, as well as other mechanical derangements which may be caused by trauma to the tissue, such as a displacement of torn meniscus tissue, meniscectomy, a laxation of a joint by a torn ligament, malignment of joints, bone fracture, or by hereditary disease. The present reparative method is also useful  
20 for remodeling cartilage matrix, such as in plastic or reconstructive surgery, as well as periodontal surgery. The present method may also be applied to improving a previous reparative procedure, for example, following surgical repair of a meniscus, ligament, or cartilage. Furthermore, it may prevent the onset or exacerbation of degenerative disease if applied early enough after trauma.

25 In one embodiment of the present invention, the subject method comprises treating the afflicted connective tissue with a therapeutically sufficient amount of a subject compound to regulate a cartilage repair response in the connective tissue by managing the rate of differentiation and/or proliferation of chondrocytes embedded in the tissue. Such connective tissues as articular cartilage, interarticular cartilage (menisci), costal cartilage,  
30 (connecting the true ribs and the sternum), ligaments, and tendons are particularly amenable to treatment in reconstructive and/or regenerative therapies using the subject

method. As used herein, regenerative therapies include treatment of degenerative states which have progressed to the point of which impairment of the tissue is obviously manifest, as well as preventive treatments of tissue where degeneration is in its earliest stages or imminent.

5 In an illustrative embodiment, the subject method can be used as part of a therapeutic intervention in the treatment of cartilage of a diarthroidal joint, such as a knee, an ankle, an elbow, a hip, a wrist, a knuckle of either a finger or toe, or a tempomandibular joint. The treatment can be directed to the meniscus of the joint, to the articular cartilage of the joint, or both. To further illustrate, the subject method can be used to treat a  
10 degenerative disorder of a knee, such as which might be the result of traumatic injury (e.g., a sports injury or excessive wear) or osteoarthritis. The subject regulators may be administered as an injection into the joint with, for instance, an arthroscopic needle. In some instances, the injected agent can be in the form of a hydrogel or other slow release vehicle described above in order to permit a more extended and regular contact of the  
15 agent with the treated tissue.

The present invention further contemplates the use of the subject method in the field of cartilage transplantation and prosthetic device therapies. However, problems arise, for instance, because the characteristics of cartilage and fibrocartilage varies between different tissue: such as between articular, meniscal cartilage, ligaments, and tendons,  
20 between the two ends of the same ligament or tendon, and between the superficial and deep parts of the tissue. The zonal arrangement of these tissues may reflect a gradual change in mechanical properties, and failure occurs when implanted tissue, which has not differentiated under those conditions, lacks the ability to appropriately respond. For instance, when meniscal cartilage is used to repair anterior cruciate ligaments, the tissue  
25 undergoes a metaplasia to pure fibrous tissue. By regulating the rate of chondrogenesis, the subject method can be used to particularly address this problem, by helping to adaptively control the implanted cells in the new environment and effectively resemble hypertrophic chondrocytes of an earlier developmental stage of the tissue.

In similar fashion, the subject method can be applied to enhancing both the  
30 generation of prosthetic cartilage devices and to their implantation. The need for improved treatment has motivated research aimed at creating new cartilage that is based on collagen-

glycosaminoglycan templates (Stone et al. (1990) *Clin Orthop Relat Res* 252:129), isolated chondrocytes (Grande et al. (1989) *J Orthop Res* 7:208; and Takigawa et al. (1987) *Bone Miner* 2:449), and chondrocytes attached to natural or synthetic polymers (Walitani et al. (1989) *J Bone Jt Surg* 71B:74; Vacanti et al. (1991) *Plast Reconstr Surg* 88:753; von Schroeder et al. (1991) *J Biomed Mater Res* 25:329; Freed et al. (1993) *J Biomed Mater Res* 27:11; and the Vacanti et al. U.S. Patent No. 5,041,138); For example, chondrocytes can be grown in culture on biodegradable, biocompatible highly porous scaffolds formed from polymers such as polyglycolic acid, polylactic acid, agarose gel, or other polymers which degrade over time as function of hydrolysis of the polymer backbone into innocuous monomers. The matrices are designed to allow adequate nutrient and gas exchange to the cells until engraftment occurs. The cells can be cultured *in vitro* until adequate cell volume and density has developed for the cells to be implanted. One advantage of the matrices is that they can be cast or molded into a desired shape on an individual basis, so that the final product closely resembles the patient's own ear or nose (by way of example), or flexible matrices can be used which allow for manipulation at the time of implantation, as in a joint.

In one embodiment of the subject method, the implants are contacted with a subject compound during certain stages of the culturing process in order to manage the rate of differentiation of chondrocytes and the formation of hypertrophic chondrocytes in the culture.

In another embodiment, the implanted device is treated with a subject compound in order to actively remodel the implanted matrix and to make it more suitable for its intended function. As set out above with respect to tissue transplants, the artificial transplants suffer from the same deficiency of not being derived in a setting which is comparable to the actual mechanical environment in which the matrix is implanted. The ability to regulate the chondrocytes in the matrix by the subject method can allow the implant to acquire characteristics similar to the tissue for which it is intended to replace.

In yet another embodiment, the subject method is used to enhance attachment of prosthetic devices. To illustrate, the subject method can be used in the implantation of a periodontal prosthesis, wherein the treatment of the surrounding connective tissue stimulates formation of periodontal ligament about the prosthesis.

In still further embodiments, the subject method can be employed as part of a regimen for the generation of bone (osteogenesis) at a site in the animal where such skeletal tissue is deficient. Indian hedgehog is particularly associated with the hypertrophic chondrocytes that are ultimately replaced by osteoblasts. For instance, administration of a compound of the present invention can be employed as part of a method for regulating the rate of bone loss in a subject. For example, preparations comprising subject compounds can be employed, for example, to control endochondral ossification in the formation of a "model" for ossification.

In yet another embodiment of the present invention, a subject compound can be used to regulate spermatogenesis. The *hedgehog* proteins, particularly *Dhh*, have been shown to be involved in the differentiation and/or proliferation and maintenance of testicular germ cells. *Dhh* expression is initiated in Sertoli cell precursors shortly after the activation of *Sry* (testicular determining gene) and persists in the testis into the adult. Males are viable but infertile, owing to a complete absence of mature sperm. Examination of the developing testis in different genetic backgrounds suggests that *Dhh* regulates both early and late stages of spermatogenesis. Bitgood et al. (1996) Curr Biol 6:298. In a preferred embodiment, the subject compound can be used as a contraceptive. In similar fashion, compounds of the subject method are potentially useful for modulating normal ovarian function.

The subject method also has wide applicability to the treatment or prophylaxis of disorders afflicting epithelial tissue, as well as in cosmetic uses. In general, the method can be characterized as including a step of administering to an animal an amount of a subject compound effective to alter the growth state of a treated epithelial tissue. The mode of administration and dosage regimens will vary depending on the epithelial tissue(s) which is to be treated. For example, topical formulations will be preferred where the treated tissue is epidermal tissue, such as dermal or mucosal tissues.

A method which "promotes the healing of a wound" results in the wound healing more quickly as a result of the treatment than a similar wound heals in the absence of the treatment. "Promotion of wound healing" can also mean that the method regulates the proliferation and/or growth of, *inter alia*, keratinocytes, or that the wound heals with less scarring, less wound contraction, less collagen deposition and more superficial surface area. In certain instances, "promotion of wound healing" can also mean that certain

methods of wound healing have improved success rates, (e.g., the take rates of skin grafts) when used together with the method of the present invention.

Despite significant progress in reconstructive surgical techniques, scarring can be an important obstacle in regaining normal function and appearance of healed skin. This is particularly true when pathologic scarring such as keloids or hypertrophic scars of the hands or face causes functional disability or physical deformity. In the severest circumstances, such scarring may precipitate psychosocial distress and a life of economic deprivation. Wound repair includes the stages of hemostasis, inflammation, proliferation, and remodeling. The proliferative stage involves multiplication of fibroblasts and endothelial and epithelial cells. Through the use of the subject method, the rate of proliferation of epithelial cells in and proximal to the wound can be controlled in order to accelerate closure of the wound and/or minimize the formation of scar tissue.

The present treatment can also be effective as part of a therapeutic regimen for treating oral and paraoral ulcers, e.g., resulting from radiation and/or chemotherapy. Such ulcers commonly develop within days after chemotherapy or radiation therapy. These ulcers usually begin as small, painful irregularly shaped lesions usually covered by a delicate gray necrotic membrane and surrounded by inflammatory tissue. In many instances, lack of treatment results in proliferation of tissue around the periphery of the lesion on an inflammatory basis. For instance, the epithelium bordering the ulcer usually demonstrates proliferative activity, resulting in loss of continuity of surface epithelium. These lesions, because of their size and loss of epithelial integrity, dispose the body to potential secondary infection. Routine ingestion of food and water becomes a very painful event and, if the ulcers proliferate throughout the alimentary canal, diarrhea usually is evident with all its complicating factors. According to the present invention, a treatment for such ulcers which includes application of a subject compound can reduce the abnormal proliferation and differentiation of the affected epithelium, helping to reduce the severity of subsequent inflammatory events.

The subject method and compositions can also be used to treat wounds resulting from dermatological diseases, such as lesions resulting from autoimmune disorders such as psoriasis. Atopic dermatitis refers to skin trauma resulting from allergies associated with an immune response caused by allergens such as pollens, foods, dander, insect venoms and plant toxins.

In other embodiments, antiproliferative preparations of subject compounds can be used to inhibit lens epithelial cell proliferation to prevent post-operative complications of extracapsular cataract extraction. Cataract is an intractable eye disease and various studies on a treatment of cataract have been made. But at present, the treatment of cataract is attained by surgical operations. Cataract surgery has been applied for a long time and various operative methods have been examined. Extracapsular lens extraction has become the method of choice for removing cataracts. The major medical advantages of this technique over intracapsular extraction are lower incidence of aphakic cystoid macular edema and retinal detachment. Extracapsular extraction is also required for implantation of posterior chamber type intraocular lenses which are now considered to be the lenses of choice in most cases.

However, a disadvantage of extracapsular cataract extraction is the high incidence of posterior lens capsule opacification, often called after-cataract, which can occur in up to 50% of cases within three years after surgery. After-cataract is caused by proliferation of equatorial and anterior capsule lens epithelial cells which remain after extracapsular lens extraction. These cells proliferate to cause Sommerling rings, and along with fibroblasts which also deposit and occur on the posterior capsule, cause opacification of the posterior capsule, which interferes with vision. Prevention of after-cataract would be preferable to treatment. To inhibit secondary cataract formation, the subject method provides a means for inhibiting proliferation of the remaining lens epithelial cells. For example, such cells can be induced to remain quiescent by instilling a solution containing a preparation of a subject compound into the anterior chamber of the eye after lens removal. Furthermore, the solution can be osmotically balanced to provide minimal effective dosage when instilled into the anterior chamber of the eye, thereby inhibiting subcapsular epithelial growth with some specificity.

The subject method can also be used in the treatment of corneopathies marked by corneal epithelial cell proliferation, as for example in ocular epithelial disorders such as epithelial downgrowth or squamous cell carcinomas of the ocular surface.

Levine et al. (1997) J Neurosci 17:6277 show that hedgehog proteins can regulate mitogenesis and photoreceptor differentiation in the vertebrate retina, and *Ihh* is a candidate factor from the pigmented epithelium to promote retinal progenitor proliferation and photoreceptor differentiation. Likewise, Jensen et al. (1997) Development 124:363

demonstrated that treatment of cultures of perinatal mouse retinal cells with the amino-terminal fragment of *Sonic hedgehog* results in an increase in the proportion of cells that incorporate bromodeoxuridine, in total cell numbers, and in rod photoreceptors, amacrine cells and Muller glial cells, suggesting that *Sonic hedgehog* promotes the proliferation of retinal precursor cells. Thus, the subject method can be used in the treatment of proliferative diseases of retinal cells and regulate photoreceptor differentiation.

Yet another aspect of the present invention relates to the use of the subject method to control hair growth. Hair is basically composed of keratin, a tough and insoluble protein; its chief strength lies in its disulphide bond of cystine. Each individual hair comprises a cylindrical shaft and a root, and is contained in a follicle, a flask-like depression in the skin. The bottom of the follicle contains a finger-like projection termed the papilla, which consists of connective tissue from which hair grows, and through which blood vessels supply the cells with nourishment. The shaft is the part that extends outwards from the skin surface, whilst the root has been described as the buried part of the hair. The base of the root expands into the hair bulb, which rests upon the papilla. Cells from which the hair is produced grow in the bulb of the follicle; they are extruded in the form of fibers as the cells proliferate in the follicle. Hair "growth" refers to the formation and elongation of the hair fiber by the dividing cells.

As is well known in the art, the common hair cycle is divided into three stages: anagen, catagen and telogen. During the active phase (anagen), the epidermal stem cells of the dermal papilla divide rapidly. Daughter cells move upward and differentiate to form the concentric layers of the hair itself. The transitional stage, catagen, is marked by the cessation of mitosis of the stem cells in the follicle. The resting stage is known as telogen, where the hair is retained within the scalp for several weeks before an emerging new hair developing below it dislodges the telogen-phase shaft from its follicle. From this model it has become clear that the larger the pool of dividing stem cells that differentiate into hair cells, the more hair growth occurs. Accordingly, methods for increasing or reducing hair growth can be carried out by potentiating or inhibiting, respectively, the proliferation of these stem cells.

In certain embodiments, the subject method can be employed as a way of reducing the growth of human hair as opposed to its conventional removal by cutting, shaving, or depilation. For instance, the present method can be used in the treatment of trichosis

characterized by abnormally rapid or dense growth of hair, e.g. hypertrichosis. In an exemplary embodiment, subject compounds can be used to manage hirsutism, a disorder marked by abnormal hairiness. The subject method can also provide a process for extending the duration of depilation.

5           Moreover, because a subject compound will often be cytostatic to epithelial cells, rather than cytotoxic, such agents can be used to protect hair follicle cells from cytotoxic agents which require progression into S-phase of the cell-cycle for efficacy, e.g. radiation-induced death. Treatment by the subject method can provide protection by causing the hair follicle cells to become quiescent, e.g., by inhibiting the cells from entering S phase, and  
10           thereby preventing the follicle cells from undergoing mitotic catastrophe or programmed cell death. For instance, subject compounds can be used for patients undergoing chemo- or radiation-therapies which ordinarily result in hair loss. By inhibiting cell-cycle progression during such therapies, the subject treatment can protect hair follicle cells from death which might otherwise result from activation of cell death programs. After the therapy has  
15           concluded, the instant method can also be removed with concomitant relief of the inhibition of follicle cell proliferation.

          The subject method can also be used in the treatment of folliculitis, such as folliculitis decalvans, folliculitis ulerythematososa reticulata or keloid folliculitis. For example, a cosmetic preparation of a subject compound can be applied topically in the  
20           treatment of pseudofolliculitis, a chronic disorder occurring most often in the submandibular region of the neck and associated with shaving, the characteristic lesions of which are erythematous papules and pustules containing buried hairs.

          In another aspect of the invention, the subject method can be used to induce differentiation and/or inhibit proliferation of epithelially derived tissue. Such forms of  
25           these molecules can provide a basis for differentiation therapy for the treatment of hyperplastic and/or neoplastic conditions involving epithelial tissue. For example, such preparations can be used for the treatment of cutaneous diseases in which there is abnormal proliferation or growth of cells of the skin.

          For instance, the pharmaceutical preparations of the invention are intended for the  
30           treatment of hyperplastic epidermal conditions, such as keratosis, as well as for the treatment of neoplastic epidermal conditions such as those characterized by a high proliferation rate for various skin cancers, as for example basal cell carcinoma or



squamous cell carcinoma. The subject method can also be used in the treatment of autoimmune diseases affecting the skin, in particular, of dermatological diseases involving morbid proliferation and/or keratinization of the epidermis, as for example, caused by psoriasis or atopic dermatosis.

5 Many common diseases of the skin, such as psoriasis, squamous cell carcinoma, keratoacanthoma and actinic keratosis are characterized by localized abnormal proliferation and growth. For example, in psoriasis, which is characterized by scaly, red, elevated plaques on the skin, the keratinocytes are known to proliferate much more rapidly than normal and to differentiate less completely.

10 In one embodiment, the preparations of the present invention are suitable for the treatment of dermatological ailments linked to keratinization disorders causing abnormal proliferation of skin cells, which disorders may be marked by either inflammatory or non-inflammatory components. To illustrate, therapeutic preparations of a subject compound, e.g., which promotes quiescence or differentiation, can be used to treat varying forms of  
15 psoriasis, be they cutaneous, mucosal or ungual. Psoriasis, as described above, is typically characterized by epidermal keratinocytes which display marked proliferative activation and differentiation along a "regenerative" pathway. Treatment with an antiproliferative embodiment of the subject method can be used to reverse the pathological epidermal activation and can provide a basis for sustained remission of the disease.

20 A variety of other keratotic lesions are also candidates for treatment with the subject method. Actinic keratoses, for example, are superficial inflammatory premalignant tumors arising on sun-exposed and irradiated skin. The lesions are erythematous to brown with variable scaling. Current therapies include excisional and cryosurgery. These treatments are painful, however, and often produce cosmetically unacceptable scarring.  
25 Accordingly, treatment of keratosis, such as actinic keratosis, can include application, preferably topical, of a subject compound composition in amounts sufficient to inhibit hyperproliferation of epidermal/epidermoid cells of the lesion.

Acne represents yet another dermatologic ailment which may be treated by the subject method. Acne vulgaris, for instance, is a multifactorial disease most commonly  
30 occurring in teenagers and young adults, and is characterized by the appearance of inflammatory and noninflammatory lesions on the face and upper trunk. The basic defect which gives rise to acne vulgaris is hypercornification of the duct of a hyperactive

sebaceous gland. Hypercornification blocks the normal mobility of skin and follicle microorganisms, and in so doing, stimulates the release of lipases by *Propionobacterium acnes* and *Staphylococcus epidermidis* bacteria and *Pitrosporum ovale*, a yeast. Treatment with an antiproliferative subject compound, particularly topical preparations, may be  
5 useful for preventing the transitional features of the ducts, e.g. hypercornification, which lead to lesion formation. The subject treatment may further include, for example, antibiotics, retinoids and antiandrogens.

The present invention also provides a method for treating various forms of dermatitis. Dermatitis is a descriptive term referring to poorly demarcated lesions which  
10 are either pruritic, erythematous, scaly, blistered, weeping, fissured or crusted. These lesions arise from any of a wide variety of causes. The most common types of dermatitis are atopic, contact and diaper dermatitis. For instance, seborrheic dermatitis is a chronic, usually pruritic, dermatitis with erythema, dry, moist, or greasy scaling, and yellow crusted patches on various areas, especially the scalp, with exfoliation of an excessive amount of  
15 dry scales. The subject method can also be used in the treatment of stasis dermatitis, an often chronic, usually eczematous dermatitis. Actinic dermatitis is dermatitis that due to exposure to actinic radiation such as that from the sun, ultraviolet waves or x- or gamma-radiation. According to the present invention, the subject method can be used in the treatment and/or prevention of certain symptoms of dermatitis caused by unwanted  
20 proliferation of epithelial cells. Such therapies for these various forms of dermatitis can also include topical and systemic corticosteroids, antipuritics, and antibiotics.

Ailments which may be treated by the subject method are disorders specific to non-humans, such as mange.

In still another embodiment, the subject method can be used in the treatment of  
25 human cancers, particularly basal cell carcinomas and other tumors of epithelial tissues such as the skin. For example, subject compounds can be employed, in the subject method, as part of a treatment for basal cell nevus syndrome (BCNS), and other human carcinomas, adenocarcinomas, sarcomas and the like.

In a preferred embodiment, the subject method is used as part of a treatment of  
30 prophylaxis regimen for treating (or preventing) basal cell carcinoma. The deregulation of the *hedgehog* signaling pathway may be a general feature of basal cell carcinomas caused by *ptc* mutations. Consistent overexpression of human *ptc* mRNA has been described in

tumors of familial and sporadic BCCs, determined by in situ hybridization. Mutations that inactivate *ptc* may be expected to result in overexpression of mutant *Ptc*, because *ptc* displays negative autoregulation. Prior research demonstrates that overexpression of *hedgehog* proteins can also lead to tumorigenesis. That *sonic hedgehog* (*Shh*) has a role in

5 tumorigenesis in the mouse has been suggested by research in which transgenic mice overexpressing *Shh* in the skin developed features of BCNS, including multiple BCC-like epidermal proliferations over the entire skin surface, after only a few days of skin development. A mutation in the *Shh* human gene from a BCC was also described; it was suggested that *Shh* or other *Hh* genes in humans could act as dominant oncogenes in

10 humans. Sporadic *ptc* mutations have also been observed in BCCs from otherwise normal individuals, some of which are UV-signature mutations. In one recent study of sporadic BCCs, five UV-signature type mutations, either CT or CCTT changes, were found out of fifteen tumors determined to contain *ptc* mutations. Another recent analysis of sporadic *ptc* mutations in BCCs and neuroectodermal tumors revealed one CT change in one of three

15 *ptc* mutations found in the BCCs. See, for example, Goodrich et al. (1997) Science 277:1109-13; Xie et al. (1997) Cancer Res 57:2369-72; Oro et al. (1997) Science 276:817-21; Xie et al. (1997) Genes Chromosomes Cancer 18:305-9; Stone et al. (1996) Nature 384:129-34; and Johnson et al. (1996) Science 272:1668-71.

The subject method can also be used to treat patients with BCNS, e.g., to prevent

20 BCC or other effects of the disease which may be the result of *ptc* loss-of-function, *hedgehog* gain-of-function, or *smoothed* gain-of-function. Basal cell nevus syndrome is a rare autosomal dominant disorder characterized by multiple BCCs that appear at a young age. BCNS patients are very susceptible to the development of these tumors; in the second decade of life, large numbers appear, mainly on sun-exposed areas of the skin. This disease

25 also causes a number of developmental abnormalities, including rib, head and face alterations, and sometimes polydactyly, syndactyly, and spina bifida. They also develop a number of tumor types in addition to BCCs: fibromas of the ovaries and heart, cysts of the skin and jaws, and in the central nervous system, medulloblastomas and meningiomas. The subject method can be used to prevent or treat such tumor types in BCNS and non-BCNS

30 patients. Studies of BCNS patients show that they have both genomic and sporadic mutations in the *ptc* gene, suggesting that these mutations are the ultimate cause of this disease.

In another aspect, the present invention provides pharmaceutical preparations and methods for controlling the formation of megakaryocyte-derived cells and/or controlling the functional performance of megakaryocyte-derived cells. For instance, certain of the compositions disclosed herein may be applied to the treatment or prevention of a variety  
5 hyperplastic or neoplastic conditions affecting platelets.

In another aspect, the present invention provides pharmaceutical preparations comprising the subject compounds. The compounds for use in the subject method may be conveniently formulated for administration with a biologically acceptable and/or sterile medium, such as water, buffered saline, polyol (for example, glycerol, propylene glycol,  
10 liquid polyethylene glycol and the like) or suitable mixtures thereof. The optimum concentration of the active ingredient(s) in the chosen medium can be determined empirically, according to procedures well known to medicinal chemists. As used herein, "biologically acceptable medium" includes any and all solvents, dispersion media, and the like which may be appropriate for the desired route of administration of the pharmaceutical  
15 preparation. The use of such media for pharmaceutically active substances is known in the art. Except insofar as any conventional media or agent is incompatible with the activity of the subject compounds, its use in the pharmaceutical preparation of the invention is contemplated. Suitable vehicles and their formulation inclusive of other proteins are described, for example, in the book *Remington's Pharmaceutical Sciences* (Remington's  
20 *Pharmaceutical Sciences*. Mack Publishing Company, Easton, Pa., USA 1985). These vehicles include injectable "deposit formulations".

Pharmaceutical formulations of the present invention can also include veterinary compositions, e.g., pharmaceutical preparations of the subject compounds suitable for veterinary uses, e.g., for the treatment of live stock or domestic animals, e.g., dogs.

25 Methods of introduction may also be provided by rechargeable or biodegradable devices. Various slow release polymeric devices have been developed and tested *in vivo* in recent years for the controlled delivery of drugs, including proteinacious biopharmaceuticals. A variety of biocompatible polymers (including hydrogels), including both biodegradable and non-degradable polymers, can be used to form an implant for the  
30 sustained release of a subject compound at a particular target site.

The preparations of the present invention may be given orally, parenterally, topically, or rectally. They are of course given by forms suitable for each administration route. For example, they are administered in tablets or capsule form, by injection, inhalation, eye lotion, ointment, suppository, controlled release patch, etc. administration  
5 by injection, infusion or inhalation; topical by lotion or ointment; and rectal by suppositories. Oral and topical administrations are preferred.

The phrases "parenteral administration" and "administered parenterally" as used herein means modes of administration other than enteral and topical administration, usually by injection, and includes, without limitation, intravenous, intramuscular,  
10 intraarterial, intrathecal, intracapsular, intraorbital, intracardiac, intradermal, intraperitoneal, transtracheal, subcutaneous, subcuticular, intraarticular, subcapsular, subarachnoid, intraspinal and intrasternal injection and infusion.

The phrases "systemic administration," "administered systemically," "peripheral administration" and "administered peripherally" as used herein mean the administration of  
15 a compound, drug or other material other than directly into the central nervous system, such that it enters the patient's system and, thus, is subject to metabolism and other like processes, for example, subcutaneous administration.

These compounds may be administered to humans and other animals for therapy by any suitable route of administration, including orally, nasally, as by, for example, a spray,  
20 rectally, intravaginally, parenterally, intracisternally and topically, as by powders, ointments or drops, including buccally and sublingually.

Regardless of the route of administration selected, the compounds of the present invention, which may be used in a suitable hydrated form, and/or the pharmaceutical compositions of the present invention, are formulated into pharmaceutically acceptable  
25 dosage forms such as described below or by other conventional methods known to those of skill in the art.

Actual dosage levels of the active ingredients in the pharmaceutical compositions of this invention may be varied so as to obtain an amount of the active ingredient which is effective to achieve the desired therapeutic response for a particular patient, composition,  
30 and mode of administration, without being toxic to the patient.

The selected dosage level will depend upon a variety of factors including the activity of the particular compound of the present invention employed, or the ester, salt or amide thereof, the route of administration, the time of administration, the rate of excretion of the particular compound being employed, the duration of the treatment, other drugs, compounds and/or materials used in combination with the particular compound employed, the age, sex, weight, condition, general health and prior medical history of the patient being treated, and like factors well known in the medical arts.

A physician or veterinarian having ordinary skill in the art can readily determine and prescribe the effective amount of the pharmaceutical composition required. For example, the physician or veterinarian could start doses of the compounds of the invention employed in the pharmaceutical composition at levels lower than that required in order to achieve the desired therapeutic effect and gradually increase the dosage until the desired effect is achieved.

In general, a suitable daily dose of a compound of the invention will be that amount of the compound which is the lowest dose effective to produce a therapeutic effect. Such an effective dose will generally depend upon the factors described above. Generally, intravenous, intracerebroventricular and subcutaneous doses of the compounds of this invention for a patient will range from about 0.0001 to about 100 mg per kilogram of body weight per day.

If desired, the effective daily dose of the active compound may be administered as two, three, four, five, six or more sub-doses administered separately at appropriate intervals throughout the day, optionally, in unit dosage forms.

The term "treatment" is intended to encompass also prophylaxis, therapy and cure.

The patient receiving this treatment is any animal in need, including primates, in particular humans, and other mammals such as equines, cattle, swine and sheep; and poultry and pets in general.

The compound of the invention can be administered as such or in admixtures with pharmaceutically acceptable and/or sterile carriers and can also be administered in conjunction with other antimicrobial agents such as penicillins, cephalosporins, aminoglycosides and glycopeptides. Conjunctive therapy, thus includes sequential, simultaneous and separate administration of the active compound in a way that the

therapeutical effects of the first administered one is not entirely disappeared when the subsequent is administered.

#### V. Pharmaceutical Compositions

5 While it is possible for a compound of the present invention to be administered alone, it is preferable to administer the compound as a pharmaceutical formulation (composition). The subject compounds according to the invention may be formulated for administration in any convenient way for use in human or veterinary medicine. In certain  
10 embodiments, the compound included in the pharmaceutical preparation may be active itself, or may be a prodrug, e.g., capable of being converted to an active compound in a physiological setting.

Thus, another aspect of the present invention provides pharmaceutically acceptable compositions comprising a therapeutically effective amount of one or more of the compounds described above, formulated together with one or more pharmaceutically  
15 acceptable carriers (additives) and/or diluents. As described in detail below, the pharmaceutical compositions of the present invention may be specially formulated for administration in solid or liquid form, including those adapted for the following: (1) oral administration, for example, drenches (aqueous or non-aqueous solutions or suspensions), tablets, boluses, powders, granules, pastes for application to the tongue; (2) parenteral  
20 administration, for example, by subcutaneous, intramuscular or intravenous injection as, for example, a sterile solution or suspension; (3) topical application, for example, as a cream, ointment or spray applied to the skin; or (4) intravaginally or intrarectally, for example, as a pessary, cream or foam. However, in certain embodiments the subject compounds may be simply dissolved or suspended in sterile water. In certain  
25 embodiments, the pharmaceutical preparation is non-pyrogenic, i.e., does not elevate the body temperature of a patient.

The phrase "therapeutically effective amount" as used herein means that amount of a compound, material, or composition comprising a compound of the present invention which is effective for producing some desired therapeutic effect, e.g., by overcoming a *ptc*  
30 loss-of-function, *hedgehog* gain-of-function, or *smoothened* gain-of-function, in at least a sub-population of cells in an animal and thereby blocking the biological consequences of

that pathway in the treated cells, at a reasonable benefit/risk ratio applicable to any medical treatment.

The phrase "pharmaceutically acceptable" is employed herein to refer to those compounds, materials, compositions, and/or dosage forms which are, within the scope of sound medical judgment, suitable for use in contact with the tissues of human beings and animals without excessive toxicity, irritation, allergic response, or other problem or complication, commensurate with a reasonable benefit/risk ratio.

The phrase "pharmaceutically acceptable carrier" as used herein means a pharmaceutically acceptable material, composition or vehicle, such as a liquid or solid filler, diluent, excipient, solvent or encapsulating material, involved in carrying or transporting the subject regulators from one organ, or portion of the body, to another organ, or portion of the body. Each carrier must be "acceptable" in the sense of being compatible with the other ingredients of the formulation and not injurious to the patient. Some examples of materials which can serve as pharmaceutically acceptable carriers include: (1) sugars, such as lactose, glucose and sucrose; (2) starches, such as corn starch and potato starch; (3) cellulose, and its derivatives, such as sodium carboxymethyl cellulose, ethyl cellulose and cellulose acetate; (4) powdered tragacanth; (5) malt; (6) gelatin; (7) talc; (8) excipients, such as cocoa butter and suppository waxes; (9) oils, such as peanut oil, cottonseed oil, safflower oil, sesame oil, olive oil, corn oil and soybean oil; (10) glycols, such as propylene glycol; (11) polyols, such as glycerin, sorbitol, mannitol and polyethylene glycol; (12) esters, such as ethyl oleate and ethyl laurate; (13) agar; (14) buffering agents, such as magnesium hydroxide and aluminum hydroxide; (15) alginic acid; (16) pyrogen-free water; (17) isotonic saline; (18) Ringer's solution; (19) ethyl alcohol; (20) phosphate buffer solutions; and (21) other non-toxic compatible substances employed in pharmaceutical formulations.

As set out above, certain embodiments of the present compounds may contain a basic functional group, such as amino or alkylamino, and are, thus, capable of forming pharmaceutically acceptable salts with pharmaceutically acceptable acids. The term "pharmaceutically acceptable salts" in this respect, refers to the relatively non-toxic, inorganic and organic acid addition salts of compounds of the present invention. These salts can be prepared *in situ* during the final isolation and purification of the compounds of the invention, or by separately reacting a purified compound of the invention in its free



base form with a suitable organic or inorganic acid, and isolating the salt thus formed. Representative salts include the hydrobromide, hydrochloride, sulfate, bisulfate, phosphate, nitrate, acetate, valerate, oleate, palmitate, stearate, laurate, benzoate, lactate, phosphate, tosylate, citrate, maleate, fumarate, succinate, tartrate, naphthylate, mesylate, glucoheptonate, lactobionate, and laurylsulphonate salts and the like. (See, for example, Berge et al. (1977) "Pharmaceutical Salts", *J. Pharm. Sci.* 66:1-19)

The pharmaceutically acceptable salts of the subject compounds include the conventional nontoxic salts or quaternary ammonium salts of the compounds, e.g., from non-toxic organic or inorganic acids. For example, such conventional nontoxic salts include those derived from inorganic acids such as hydrochloride, hydrobromic, sulfuric, sulfamic, phosphoric, nitric, and the like; and the salts prepared from organic acids such as acetic, propionic, succinic, glycolic, stearic, lactic, malic, tartaric, citric, ascorbic, palmitic, maleic, hydroxymaleic, phenylacetic, glutamic, benzoic, salicylic, sulfanilic, 2-acetoxybenzoic, fumaric, toluenesulfonic, methanesulfonic, ethane disulfonic, oxalic, isothionic, and the like.

In other cases, the compounds of the present invention may contain one or more acidic functional groups and, thus, are capable of forming pharmaceutically acceptable salts with pharmaceutically acceptable bases. The term "pharmaceutically acceptable salts" in these instances refers to the relatively non-toxic, inorganic and organic base addition salts of compounds of the present invention. These salts can likewise be prepared *in situ* during the final isolation and purification of the compounds, or by separately reacting the purified compound in its free acid form with a suitable base, such as the hydroxide, carbonate or bicarbonate of a pharmaceutically acceptable metal cation, with ammonia, or with a pharmaceutically acceptable organic primary, secondary or tertiary amine. Representative alkali or alkaline earth salts include the lithium, sodium, potassium, calcium, magnesium, and aluminum salts and the like. Representative organic amines useful for the formation of base addition salts include ethylamine, diethylamine, ethylenediamine, ethanolamine, diethanolamine, piperazine and the like. (See, for example, Berge et al., *supra*)

Wetting agents, emulsifiers and lubricants, such as sodium lauryl sulfate and magnesium stearate, as well as coloring agents, release agents, coating agents, sweetening,

flavoring and perfuming agents, preservatives and antioxidants can also be present in the compositions.

Examples of pharmaceutically acceptable antioxidants include: (1) water soluble antioxidants, such as ascorbic acid, cysteine hydrochloride, sodium bisulfate, sodium metabisulfite, sodium sulfite and the like; (2) oil-soluble antioxidants, such as ascorbyl palmitate, butylated hydroxyanisole (BHA), butylated hydroxytoluene (BHT), lecithin, propyl gallate, alpha-tocopherol, and the like; and (3) metal chelating agents, such as citric acid, ethylenediamine tetraacetic acid (EDTA), sorbitol, tartaric acid, phosphoric acid, and the like.

Formulations of the present invention include those suitable for oral, nasal, topical (including buccal and sublingual), rectal, vaginal and/or parenteral administration. The formulations may conveniently be presented in unit dosage form and may be prepared by any methods well known in the art of pharmacy. The amount of active ingredient which can be combined with a carrier material to produce a single dosage form will vary depending upon the host being treated, the particular mode of administration. The amount of active ingredient which can be combined with a carrier material to produce a single dosage form will generally be that amount of the compound which produces a therapeutic effect. Generally, out of one hundred per cent, this amount will range from about 1 per cent to about ninety-nine percent of active ingredient, preferably from about 5 per cent to about 70 per cent, most preferably from about 10 per cent to about 30 per cent.

Methods of preparing these formulations or compositions include the step of bringing into association a compound of the present invention with the carrier and, optionally, one or more accessory ingredients. In general, the formulations are prepared by uniformly and intimately bringing into association a compound of the present invention with liquid carriers, or finely divided solid carriers, or both, and then, if necessary, shaping the product.

Formulations of the invention suitable for oral administration may be in the form of capsules, cachets, pills, tablets, lozenges (using a flavored basis, usually sucrose and acacia or tragacanth), powders, granules, or as a solution or a suspension in an aqueous or non-aqueous liquid, or as an oil-in-water or water-in-oil liquid emulsion, or as an elixir or syrup, or as pastilles (using an inert base, such as gelatin and glycerin, or sucrose and acacia) and/or as mouth washes and the like, each containing a predetermined amount of a

compound of the present invention as an active ingredient. A compound of the present invention may also be administered as a bolus, electuary or paste.

In solid dosage forms of the invention for oral administration (capsules, tablets, pills, dragees, powders, granules and the like), the active ingredient is mixed with one or more pharmaceutically acceptable carriers, such as sodium citrate or dicalcium phosphate, and/or any of the following: (1) fillers or extenders, such as starches, lactose, sucrose, glucose, mannitol, and/or silicic acid; (2) binders, such as, for example, carboxymethylcellulose, alginates, gelatin, polyvinyl pyrrolidone, sucrose and/or acacia; (3) humectants, such as glycerol; (4) disintegrating agents, such as agar-agar, calcium carbonate, potato or tapioca starch, alginic acid, certain silicates, and sodium carbonate; (5) solution retarding agents, such as paraffin; (6) absorption accelerators, such as quaternary ammonium compounds; (7) wetting agents, such as, for example, cetyl alcohol and glycerol monostearate; (8) absorbents, such as kaolin and bentonite clay; (9) lubricants, such as talc, calcium stearate, magnesium stearate, solid polyethylene glycols, sodium lauryl sulfate, and mixtures thereof; and (10) coloring agents. In the case of capsules, tablets and pills, the pharmaceutical compositions may also comprise buffering agents. Solid compositions of a similar type may also be employed as fillers in soft and hard-filled gelatin capsules using such excipients as lactose or milk sugars, as well as high molecular weight polyethylene glycols and the like.

A tablet may be made by compression or molding, optionally with one or more accessory ingredients. Compressed tablets may be prepared using binder (for example, gelatin or hydroxypropylmethyl cellulose), lubricant, inert diluent, preservative, disintegrant (for example, sodium starch glycolate or cross-linked sodium carboxymethyl cellulose), surface-active or dispersing agent. Molded tablets may be made by molding in a suitable machine a mixture of the powdered compound moistened with an inert liquid diluent.

The tablets, and other solid dosage forms of the pharmaceutical compositions of the present invention, such as dragees, capsules, pills and granules, may optionally be scored or prepared with coatings and shells, such as enteric coatings and other coatings well known in the pharmaceutical-formulating art. They may also be formulated so as to provide slow or controlled release of the active ingredient therein using, for example, hydroxypropylmethyl cellulose in varying proportions to provide the desired release

profile, other polymer matrices, liposomes and/or microspheres. They may be sterilized by, for example, filtration through a bacteria-retaining filter, or by incorporating sterilizing agents in the form of sterile solid compositions which can be dissolved in sterile water, or some other sterile injectable medium immediately before use. These compositions may also optionally contain opacifying agents and may be of a composition that they release the active ingredient(s) only, or preferentially, in a certain portion of the gastrointestinal tract, optionally, in a delayed manner. Examples of embedding compositions which can be used include polymeric substances and waxes. The active ingredient can also be in micro-encapsulated form, if appropriate, with one or more of the above-described excipients.

Liquid dosage forms for oral administration of the compounds of the invention include pharmaceutically acceptable emulsions, microemulsions, solutions, suspensions, syrups and elixirs. In addition to the active ingredient, the liquid dosage forms may contain inert diluents commonly used in the art, such as, for example, water or other solvents, solubilizing agents and emulsifiers, such as ethyl alcohol, isopropyl alcohol, ethyl carbonate, ethyl acetate, benzyl alcohol, benzyl benzoate, propylene glycol, 1,3-butylene glycol, oils (in particular, cottonseed, groundnut, corn, germ, olive, castor and sesame oils), glycerol, tetrahydrofuryl alcohol, polyethylene glycols and fatty acid esters of sorbitan, and mixtures thereof.

Besides inert diluents, the oral compositions can also include adjuvants such as wetting agents, emulsifying and suspending agents, sweetening, flavoring, coloring, perfuming and preservative agents.

Suspensions, in addition to the active compounds, may contain suspending agents as, for example, ethoxylated isostearyl alcohols, polyoxyethylene sorbitol and sorbitan esters, microcrystalline cellulose, aluminum metahydroxide, bentonite, agar-agar and tragacanth, and mixtures thereof.

It is known that sterols, such as cholesterol, will form complexes with cyclodextrins. Thus, in preferred embodiments, where the inhibitor is a steroidal alkaloid, it may be formulated with cyclodextrins, such as  $\alpha$ -,  $\beta$ - and  $\gamma$ -cyclodextrin, dimethyl- $\beta$  cyclodextrin and 2-hydroxypropyl- $\beta$ -cyclodextrin.

Formulations of the pharmaceutical compositions of the invention for rectal or vaginal administration may be presented as a suppository, which may be prepared by

mixing one or more compounds of the invention with one or more suitable nonirritating excipients or carriers comprising, for example, cocoa butter, polyethylene glycol, a suppository wax or a salicylate, and which is solid at room temperature, but liquid at body temperature and, therefore, will melt in the rectum or vaginal cavity and release the active  
5 compound.

Formulations of the present invention which are suitable for vaginal administration also include pessaries, tampons, creams, gels, pastes, foams or spray formulations containing such carriers as are known in the art to be appropriate.

Dosage forms for the topical or transdermal administration of a compound of this  
10 invention include powders, sprays, ointments, pastes, creams, lotions, gels, solutions, patches and inhalants. The active compound may be mixed under sterile conditions with a pharmaceutically acceptable carrier, and with any preservatives, buffers, or propellants which may be required.

The ointments, pastes, creams and gels may contain, in addition to an active  
15 compound of this invention, excipients, such as animal and vegetable fats, oils, waxes, paraffins, starch, tragacanth, cellulose derivatives, polyethylene glycols, silicones, bentonites, silicic acid, talc and zinc oxide, or mixtures thereof.

Powders and sprays can contain, in addition to a compound of this invention, excipients such as lactose, talc, silicic acid, aluminum hydroxide, calcium silicates and  
20 polyamide powder, or mixtures of these substances. Sprays can additionally contain customary propellants, such as chlorofluorohydrocarbons and volatile unsubstituted hydrocarbons, such as butane and propane.

Transdermal patches have the added advantage of providing controlled delivery of a compound of the present invention to the body. Such dosage forms can be made by  
25 dissolving or dispersing the subject compounds in the proper medium. Absorption enhancers can also be used to increase the flux of the compound across the skin. The rate of such flux can be controlled by either providing a rate controlling membrane or dispersing the compound in a polymer matrix or gel.

Ophthalmic formulations, eye ointments, powders, solutions and the like, are also  
30 contemplated as being within the scope of this invention.

Pharmaceutical compositions of this invention suitable for parenteral administration comprise one or more compounds of the invention in combination with one or more pharmaceutically acceptable sterile isotonic aqueous or nonaqueous solutions, dispersions, suspensions or emulsions, or sterile powders which may be reconstituted into sterile injectable solutions or dispersions just prior to use, which may contain antioxidants, buffers, bacteriostats, solutes which render the formulation isotonic with the blood of the intended recipient or suspending or thickening agents.

Examples of suitable aqueous and nonaqueous carriers which may be employed in the pharmaceutical compositions of the invention include water, ethanol, polyols (such as glycerol, propylene glycol, polyethylene glycol, and the like), and suitable mixtures thereof, vegetable oils, such as olive oil, and injectable organic esters, such as ethyl oleate. Proper fluidity can be maintained, for example, by the use of coating materials, such as lecithin, by the maintenance of the required particle size in the case of dispersions, and by the use of surfactants.

These compositions may also contain adjuvants such as preservatives, wetting agents, emulsifying agents and dispersing agents. Prevention of the action of microorganisms may be ensured by the inclusion of various antibacterial and antifungal agents, for example, paraben, chlorobutanol, phenol sorbic acid, and the like. It may also be desirable to include isotonic agents, such as sugars, sodium chloride, and the like into the compositions. In addition, prolonged absorption of the injectable pharmaceutical form may be brought about by the inclusion of agents which delay absorption such as aluminum monostearate and gelatin.

In some cases, in order to prolong the effect of a drug, it is desirable to slow the absorption of the drug from subcutaneous or intramuscular injection. This may be accomplished by the use of a liquid suspension of crystalline or amorphous material having poor water solubility. The rate of absorption of the drug then depends upon its rate of dissolution which, in turn, may depend upon crystal size and crystalline form. Alternatively, delayed absorption of a parenterally administered drug form is accomplished by dissolving or suspending the drug in an oil vehicle.

Injectable depot forms are made by forming microencapsule matrices of the subject compounds in biodegradable polymers such as polylactide-polyglycolide. Depending on the ratio of drug to polymer, and the nature of the particular polymer employed, the rate of

drug release can be controlled. Examples of other biodegradable polymers include poly(orthoesters) and poly(anhydrides). Depot injectable formulations are also prepared by entrapping the drug in liposomes or microemulsions which are compatible with body tissue.

- 5           When the compounds of the present invention are administered as pharmaceuticals, to humans and animals, they can be given per se or as a pharmaceutical composition containing, for example, 0.1 to 99.5% (more preferably, 0.5 to 90%) of active ingredient in combination with a pharmaceutically acceptable carrier.

- 10           The addition of the active compound of the invention to animal feed is preferably accomplished by preparing an appropriate feed premix containing the active compound in an effective amount and incorporating the premix into the complete ration.

- 15           Alternatively, an intermediate concentrate or feed supplement containing the active ingredient can be blended into the feed. The way in which such feed premixes and complete rations can be prepared and administered are described in reference books (such as "Applied Animal Nutrition", W.H. Freedman and CO., San Francisco, U.S.A., 1969 or "Livestock Feeds and Feeding" O and B books, Corvallis, Ore., U.S.A., 1977).

#### VI. Synthetic Schemes and Identification of Active Regulators

- 20           The subject compounds, and derivatives thereof, can be prepared readily by employing known synthetic methodology. As is well known in the art, these coupling reactions are carried out under relatively mild conditions and tolerate a wide range of "spectator" functionality. Additional compounds may be synthesized and tested in a combinatorial fashion, to facilitate the identification of additional compounds which may be employed in the subject method.

25

##### *a. Combinatorial Libraries*

- 30           The compounds of the present invention, particularly libraries of variants having various representative classes of substituents, are amenable to combinatorial chemistry and other parallel synthesis schemes (see, for example, PCT WO 94/08051). The result is that large libraries of related compounds, e.g. a variegated library of compounds represented

above, can be screened rapidly in high throughput assays in order to identify potential *hedgehog* regulator lead compounds, as well as to refine the specificity, toxicity, and/or cytotoxic-kinetic profile of a lead compound. For instance, *ptc*, *hedgehog*, or *smoothened* bioactivity assays, such as may be developed using cells with either a *ptc* loss-of-function, 5 *hedgehog* gain-of-function, or *smoothened* gain-of-function, can be used to screen a library of the subject compounds for those having agonist activity toward *ptc* or antagonist activity towards *hedgehog* or *smoothened*. Alternatively, bioactivity assays using cells with either a *ptc* gain-of-function, *hedgehog* loss-of-function, or *smoothened* loss-of-function, can be used to screen a library of the subject compounds for those having antagonist 10 activity toward *ptc* or agonist activity towards *hedgehog* or *smoothened*.

Simply for illustration, a combinatorial library for the purposes of the present invention is a mixture of chemically related compounds which may be screened together for a desired property. The preparation of many related compounds in a single reaction 15 greatly reduces and simplifies the number of screening processes which need to be carried out. Screening for the appropriate physical properties can be done by conventional methods.

Diversity in the library can be created at a variety of different levels. For instance, the substrate aryl groups used in the combinatorial reactions can be diverse in terms of the 20 core aryl moiety, e.g., a variegation in terms of the ring structure, and/or can be varied with respect to the other substituents.

A variety of techniques are available in the art for generating combinatorial libraries of small organic molecules such as the subject compounds. See, for example, Blondelle et al. (1995) Trends Anal. Chem. 14:83; the Affymax U.S. Patents 5,359,115 25 and 5,362,899; the Ellman U.S. Patent 5,288,514; the Still et al. PCT publication WO 94/08051; the ArQule U.S. Patents 5,736,412 and 5,712,171; Chen et al. (1994) JACS 116:2661; Kerr et al. (1993) JACS 115:252; PCT publications WO92/10092, WO93/09668 and WO91/07087; and the Lerner et al. PCT publication WO93/20242). Accordingly, a variety of libraries on the order of about 100 to 1,000,000 or more, 30 diversomers of the subject compounds can be synthesized and screened for particular activity or property.



In an exemplary embodiment, a library of candidate compound diversomers can be synthesized utilizing a scheme adapted to the techniques described in the Still et al. PCT publication WO 94/08051, e.g., being linked to a polymer bead by a hydrolyzable or photolyzable group, optionally located at one of the positions of the candidate regulators or a substituent of a synthetic intermediate. According to the Still et al. technique, the library is synthesized on a set of beads, each bead including a set of tags identifying the particular diversomer on that bead. The bead library can then be "plated" with, for example, *ptc* loss-of-function, *hedgehog* gain-of-function, or *smoothened* gain-of-function cells for which a *hedgehog* antagonist is sought. The diversomers can be released from the bead, e.g. by hydrolysis.

Many variations on the above and related pathways permit the synthesis of widely diverse libraries of compounds which may be tested as regulators of *hedgehog* function.

#### b. Screening Assays

There are a variety of assays available for determining the ability of a compound such as a *hedgehog* regulator to regulate *ptc*, *smoothened*, or *hedgehog* function, many of which can be disposed in high-throughput formats. In many drug screening programs which test libraries of compounds and natural extracts, high throughput assays are desirable in order to maximize the number of compounds surveyed in a given period of time. Thus, libraries of synthetic and natural products can be sampled for other compounds which are *hedgehog* regulators.

In addition to cell-free assays, test compounds can also be tested in cell-based assays. In one embodiment, cells which have a *ptc* loss-of-function, *hedgehog* gain-of-function, or *smoothened* gain-of-function phenotype can be contacted with a test agent of interest, with the assay scoring for, e.g., inhibition of proliferation of the cell in the presence of the test agent.

A number of gene products have been implicated in *patched*-mediated signal transduction, including *patched*, transcription factors of the *cubitus interruptus* (*ci*) family, the serine/threonine kinase *fused* (*fu*) and the gene products of *costal-2*, *smoothened* and *suppressor of fused*.

The induction of cells by hedgehog proteins sets in motion a cascade involving the activation and inhibition of downstream effectors, the ultimate consequence of which is, in some instances, a detectable change in the transcription or translation of a gene. Potential transcriptional targets of *hedgehog*-mediated signaling are the *patched* gene (Hidalgo and Ingham, 1990 Development 110, 291-301; Marigo et al., 1996 ) and the vertebrate homologs of the drosophila cubitus interruptus gene, the *GLI* genes (Hui et al. (1994) Dev Biol 162:402-413). *Patched* gene expression has been shown to be induced in cells of the limb bud and the neural plate that are responsive to *Shh*. (Marigo et al. (1996) PNAS 93:9346-51; Marigo et al. (1996) Development 122:1225-1233). The *Gli* genes encode putative transcription factors having zinc finger DNA binding domains (Orenic et al. (1990) Genes & Dev 4:1053-1067; Kinzler et al. (1990) Mol Cell Biol 10:634-642). Transcription of the *Gli* gene has been reported to be upregulated in response to *hedgehog* in limb buds, while transcription of the *Gli3* gene is downregulated in response to *hedgehog* induction (Marigo et al. (1996) Development 122:1225-1233). By selecting transcriptional regulatory sequences from such target genes, e.g., from *patched* or *Gli* genes, that are responsible for the up- or down-regulation of these genes in response to *hedgehog* signalling, and operatively linking such promoters to a reporter gene, one can derive a transcription based assay which is sensitive to the ability of a specific test compound to modify *hedgehog*-mediated signalling pathways. Expression of the reporter gene, thus, provides a valuable screening tool for the development of compounds that act as regulators of *hedgehog*.

Reporter gene based assays of this invention measure the end stage of the above described cascade of events, e.g., transcriptional modulation. Accordingly, in practicing one embodiment of the assay, a reporter gene construct is inserted into the reagent cell in order to generate a detection signal dependent on *ptc* loss-of-function, *hedgehog* gain-of-function, *smoothed* gain-of-function, or stimulation by *Shh* itself. The amount of transcription from the reporter gene may be measured using any method known to those of skill in the art to be suitable. For example, mRNA expression from the reporter gene may be detected using RNase protection or RNA-based PCR, or the protein product of the reporter gene may be identified by a characteristic stain or an intrinsic biological activity. The amount of expression from the reporter gene is then compared to the amount of expression in either the same cell in the absence of the test compound or it may be

compared with the amount of transcription in a substantially identical cell that lacks the target receptor protein. Any statistically or otherwise significant decrease in the amount of transcription indicates that the test compound has in some manner agonized the normal *ptc* signal (or antagonized the gain-of-function *hedgehog* or *smoothened* signal), e.g., the test  
5 compound is a potential *hedgehog* antagonist.

### Exemplification

The invention now being generally described, it will be more readily understood by  
10 reference to the following examples which are included merely for purposes of illustration of certain aspects and embodiments of the present invention, and are not intended to limit the invention.

#### Example 1: Steroidal Alkaloids can disrupt Hedgehog Signaling

15 In order to demonstrate an effect on Shh signaling, we chose the chick (38) as a more tractable experimental system than the rodents, sheep and other mammals in which teratogen-induced HPE predominantly has been studied (14, 15, 16, 29, 39). Chick embryos are easily cultured and manipulated and, as seen in Fig. 2, exposure of these embryos to jervine at the intermediate to definitive streak state (40) induced external  
20 malformations characteristic of HPE (similar results were obtained with cyclopamine; data not shown). The severity of these defects varied among treated embryos, as seen in panels B-E by the degree of loss of midline structures and approximation of paired lateral structures. These midline deficits thus result in the fusion of the mandibular and maxillary processes as well as the optic vesicles and olfactory processes, with consequent cyclopia  
25 and formation of a proboscis-like structure consisting of fused nasal chambers in the most severely affected embryos (Fig. 2E).

As seen in Fig. 2 in ovo treatment produced variable defects and some embryos displayed normal morphology, even at the highest concentrations tested (50μM, jervine, 5/10 and cyclopamine 2/10, data now shown). The variability of these effects may be due  
30 to imprecise embryonic staging and difficulties in applying these hydrophobic compounds

uniformly. To reduce this variability and better evaluate the potential effects of teratogenic compounds on *Shh* signaling we established an explant assay that allowed for precise tissue staging and more uniform application of the teratogen (41). As shown in Fig. 3A, medial neural plate with notochord was explanted from a region just rostral to Hensen's node. At this level, the medial neural plate does not yet express floor plate cell (HNF3 $\beta$ ) or motor neuron (Isl-1) markers (42, 43, data not shown), although the notochord does express Shh (44, 45, data not shown). As seen in Fig. 3B, after a 40 hour incubation the neural plate expresses HNF3 $\beta$  and Isl-1. Expression of these markers has been shown to depend upon *Shh* signaling, both in vivo and in vitro (2, 45), and these midline explants thus constitute an integrated assay of *Shh* signaling, comprising both inducing and target tissues.

To determine whether synthetic and plant-derived teratogens block *Shh* signaling we exposed midline explants to varying concentrations of the drugs AY 9944 and triparanol and to the steroidal alkaloids cyclopamine and jervine. As can be seen in Fig. 3D-K, all of these compounds affect *Shh* signaling, with a complete loss of HNF3 and Isl-1 expression consistently caused by sufficiently high concentrations (Fig. 3E,G,J,K). At concentrations several-fold below those required for complete inhibition, all of the teratogenic compounds are able to block HNF3 $\beta$  expression while retaining and often enhancing Isl-1 expression (Fig. 3D,F,H,I). These effects are fully consistent with inhibition of *Shh* signaling (see below). In contrast, the structurally related but not teratogenic steroidal alkaloid tomatidine (see Fig. 1, ref. 46, data not shown) is unable to block expression of HNF3 $\beta$  and Isl-1, even at concentrations two orders of magnitude higher than the inhibitory concentrations of jervine and cyclopamine (Fig. 3C).

## 25 Inhibitory compounds do not block Shh processing

Because the midline explants contain both inducing and responding tissues, we set out to distinguish possible effects of these inhibitory compounds on signal production versus possible effects on signal response. The Shh protein undergoes an intramolecular processing reaction that involves internal cleavage and gives rise to an amino-terminal product (Shh-Np) responsible for all known signaling activities. The first step of the autoprocessing reaction, mediated by the carboxy-terminal sequences within the precursor,

entails an internal rearrangement at the site of cleavage to replace the scissile peptide bond by the thioester involving a Cys side chain. In the second step cholesterol supplies the nucleophile (the  $3\beta$ -OH) that attacks the thioester intermediate, and remains covalently attached as an adduct to Shh-N<sub>p</sub> (11, 13). Autoprocessing thus is required to release active  
5 signal and the cholesterol adduct restricts the tissue distribution of the signal by causing it to associate with the cell surface (12,13).

Given this role of cholesterol in the biogenesis of Hedgehog proteins, an effect of these compounds on Shh-N<sub>p</sub> production is a particularly appealing possibility since jervine and cyclopamine structurally resemble cholesterol (Fig. 1) and AY 9944 and triparanol  
10 inhibit specific late-acting cholesterol biosynthetic enzymes (17, 18, 19, 22). To examine potential effects of these compounds on Shh processing we utilized HK293 cells cultured in lipid-depleted serum and carrying a stable integrated construct for expression of Shh under ecdysone-inducible control (47). Shh protein expression in these cells can be induced by addition of muristerone A, an ecdysone analog. As observed in embryos this  
15 protein is efficiently processed (Fig 4A, lanes 1 and 2), with little or no detectable accumulation of precursor ( $M_r$  45 kD). Addition of jervine, cyclopamine, tomatidine, AY 9944, or triparanol during the 24-hour induction period did not diminish Shh-N<sub>p</sub> production nor induce accumulation of unprocessed precursor, even at doses 5-fold higher than those required to completely inhibit Shh signaling (Figure 4A, lanes 4-13). All of the  
20 amino-terminal cleavage product generated in the presence of these compounds is detected in cell lysates, not the culture medium (data not shown), and has the same electrophoretic mobility as cholesterol-modified Shh-N<sub>p</sub>. These observations are consistent with the presence of a sterol adduct in the amino-terminal cleavage product, since lack of such an adduct is associated with release into the medium and with decreased electrophoretic  
25 mobility (the unprocessed amino-terminal fragment is designated Shh-N to distinguish it from processed Shh-N<sub>p</sub>; see lanes 8,9,17). We also failed to observe any change in efficiency of Shh processing or behavior of Shh-NP in transiently-transfected COS-7 or QT6 CELLS treated with these compounds (48). We also have observed that chick embryos treated with jervine after floor plate induction displayed the normal apical  
30 localization of Shh protein within floor plate cells (49).

Because of their structural similarity to cholesterol, we also investigated the potential effects of the plant compounds on an in vitro autoprocessing reaction utilizing

purified components. The protein utilized in this reaction is derived by replacement of all but six codons of the *Drosophila Shh* amino-terminal coding region with sequences encoding a hexahistidine purification tag (10). The resulting 29kDa protein, His6Hh-C, in purified form undergoes autoprocessing in a cholesterol-dependent manner to yield a 25kD product (50). As seen in Fig. 5A neither jervine, cyclopamine, nor tomatidine inhibit this cholesterol-stimulated autoprocessing reaction, even at concentrations 27-fold higher than that of cholesterol. Given the presence of 3 $\beta$ -OH in each of the plant compounds (Fig. 1), we also tested their ability to replace cholesterol in providing the nucleophilic group during processing. As seen in Fig. 5B, no appreciable cleavage is stimulated by addition of these compounds in the absence of cholesterol.

The observation that cholesterol synthesis inhibitors such as AY 9944 and triparanol do not inhibit processing raises the possibility that cholesterol biosynthetic precursors, which accumulate in treated cells (see below), may participate in the reaction. Figure 5C shows that the in vitro reaction can be driven by desmosterol, 7-dehydrocholesterol (7DHC), and lathosterol with efficiencies similar to that of cholesterol. Desmosterol and 7DHC are the major precursors reported to accumulate in cells treated with triparanol and AY 9944, respectively. Lanosterol, a 30 carbon cholesterol precursor, on the other hand is unable to participate in the reaction, perhaps due to steric interference by the two methyl groups attached to the C4 carbon near the 3-hydroxyl. In other studies of this in vitro reaction we have observed a requirement for an unhindered hydroxyl at the 3 $\beta$  position on a sterol nucleus, although neither the 8-carbon side chain nor the number or position(s) of the double bond(s) in the sterol nucleus appear to critically affect efficiency (51). These observations suggest that all 27 carbon sterol intermediates in the biosynthetic pathway are potential adducts in the autoprocessing reaction, and may account for the unimpaired efficiency of processing in the presence of distal synthesis inhibitors. Thus, although the extent of Shh processing in cultured cells and its localization in vivo appears to be unaffected by these inhibitory compounds (Fig. 4), we can not rule out the possibility that the sterol adduct may differ and that such an abnormally modified signal may have distinct biological properties.

**Inhibitory compounds specifically affect the response to *Shh* signaling**

Since our studies of processing provided n evidence for an inhibitory effect of these compounds on Shh signal production, we examined the alternative possibility that these compound affect response of target tissues. For these studies we utilized an intermediate neural plate explant lacking any endogenous source of inducing signal (41, see Fig. 6A).  
 5 Recombinant Shh-N protein (45, 52, 53, 54), lacking a sterol adduct, suppresses molecular markers such as Pax7 (55, see Fig. 6B, C), normally expressed in dorsal cell types, and induces ventral markers such as Isl-1 and HNF3 $\beta$  (Fig. 6D,E), normally expressed in motor neurons and floor plate cells. These cellular responses are elicited in a concentration-dependent manner, with repression of Pax7 observed at concentrations of Shh-N that are  
 10 insufficient for induction of HNF38 (ref. 55, 2 nM, Fig. 6B,C). Isl-1 and HNF3 $\beta$  occurring at the expense of Isl-1 (note that the induction of Isl-1 at 6.25 nM Shh-N in Fig. 6D is abolished at 25nM in 6E).

The teratogenic compounds are able to block completely the repression of Pax7 (at 2 nM Shh-N; Fig. 6F-I) and the induction of Isl-1 and HNF3 $\beta$  (at 25 nM Shh-N; Fig.6-S).  
 15 In addition tomatidine produces partial inhibition, but only at concentrations 100200 fold higher than those required for complete inhibition by jervine and cyclopamine (Fig. 6T). A complete inhibition of the 24 nM response to Shh-N requires doses of teratogenic compounds 2-4 higher than those required to completely block the 2 nM response; inhibition of responses to higher concentrations of Shh-N requires higher drug  
 20 concentrations. Another dose dependent effect can be noted in Fig. 6K-N, where drug concentrations two fold below the thresholds required for complete inhibition of the 25 nM response (induction of HNF3 $\beta$ ) result in retention or expansion of Isl-1 expression. A similar expansion of Isl-1 at intermediate drug concentrations was seen for midline explants (Fig. 3D-G), indicating that at a fixed level of stimulation by Shh-N, distinct  
 25 degrees of pathway activation can be produced by distinct inhibitor concentrations.

To further examine the specificity of these compounds we tested their effects on induction of a neural crest-like phenotype by BMP7. The BMP7 signaling protein is expressed in ectodermal cells adjacent to the neural plate, and appears to function in induction of neural crest and dorsal neural tube cell fates (956). To avoid contamination  
 30 with endogenous lateral signals, the explants used for these studies were taken from the ventral neural plate, but excluded the notochord and the midline (Fig 7A). Addition of BMP7 protein induced formation of migratory cells that express the HNK-1 surface

antigen (compare Figs. 7B,C), features characteristic of neural crest cells (56). Neither cell migration nor expression HNK-1 were blocked by addition of jervine at 10  $\mu$ M (Fig. 7D), a concentration exceeding that required for a complete block of Shh-N signaling. Similar results were obtained with tomatidine and with cyclopamine. These compounds also failed to inhibit formation of migratory HNK-1 positive cells from explants containing dorsal neural plate and contiguous epidermal ectoderm (49), which serves as an endogenous source of BMP activity (56).

#### Drug effects upon cholesterol homeostasis

Pervious reports indicate that triparanol and AY 944 cause the accumulation of cholesterol precursors (predominantly desmosterol and 7-dehycholesterol (7DHC) by specifically inhibiting late-acting enzymes of cholesterol biosynthesis (desmosterol  $\Delta$ 24-reductase and 7DHC  $\Delta$ 7-reductase, respectively, 17, 18, 19, 22), and a preliminary analysis of jervine also revealed an effect upon cholesterol biosynthesis (30). A direct comparison of the effects of these compounds on human primary lymphoblast cultures (57) revealed that all of them, including tomatidine, cause a relative decrease in cholesterol levels and an increase in the levels of other sterols (Table I, ref 58).

**Table 1.** Teratogenic compounds disrupt cholesterol homeostasis in cultured cells. Cholesterol biosynthesis is inhibited in primary human lymphoblasts cultured in the presence of the teratogenic compounds and tomatidine (58). The sterol profiles (57) from these cultures reveal the accumulation of multiple 27-, 28- and 29-carbon sterol precursors of cholesterol (59, 60). Esterification of PM-labeled [3H]-cholesterol in rat hepatoma cells is also inhibited by all of the compounds (63).



{Insert Table I here}

The accumulating sterols largely comprise established intermediates in the cholesterol biosynthetic pathway or closely related species that might be generated by action of the biosynthetic enzymes upon these intermediates (59). Tomatidine would appear to be the exception to this general rule, with accumulation to relatively high levels of several unusual sterols (60).

Reduction of cholesterol levels coupled with an accumulation of cholesterol biosynthetic precursors are effects observed for a group of compounds that have been termed class 2 inhibitors of cholesterol biosynthesis (61, 62). These compounds appear to act by inhibiting sterol flux between the plasma membrane (PM) and the endoplasmic reticulum (ER). Since cholesterol biosynthetic enzymes are located in the ER, and sterol precursors of cholesterol are highly concentrated in the PM, such a block in transport results in an overall reduction of cholesterol levels. We measured the effects of the synthetic and plant compounds upon esterification of exogenously added <sup>3</sup>H-labelled cholesterol (63), a process which requires transport of PM cholesterol to the ER. We observed inhibition of esterification at levels ranging from 25-75% for these compounds. An effect of AY 9944 on sterol transport previously has been reported (23), but this is the least active of the compounds we tested in inhibition of esterification. Our data therefore suggest that transport inhibition may be a factor in the effects of all of these compounds on sterol profiles, consistent with the general accumulation of multiple cholesterol biosynthetic precursors. In addition, however, AY 9944 and triparanol cause accumulation to high levels of 7DHC and desmosterol, respectively, consistent with the well-known effects of these compounds on the 7DHC  $\Delta^7$ -reductase and desmosterol  $\Delta^{24}$ -reductase enzymes.

## Discussion

The teratogenic effects of distal inhibitors of cholesterol biosynthesis have been known and studied for more than thirty years (14, 15). Similarly, cyclopamine and jervine were identified about thirty years ago as the plant compounds responsible for the teratogenic effects of the range plant *Veratrum californicum* (28, 29). The most dramatic teratogenic effect of these compounds is the induction of cyclopia and other features of

severe holoprosencephaly (HPE); the recent discovery that HPE is also caused by mutations at the murine and human loci suggested the possibility that these compounds may act to block the *Shh* signaling pathway. Our studies have verified the HPE-inducing properties of these compounds in chick embryos. We have further examined the early molecular correlates of these teratogenic effects and have demonstrated that these compounds block the induction by Shh protein of ventral cell types in chick neural plate explants.

Despite the inhibitory effects of these teratogens on cholesterol biosynthesis (17, 18, 19, 22, 30, see above), we found that none of the compounds appears to interfere with Shh processing in cultured cells, and that the plant alkaloids neither participate in nor inhibit an in vitro Hh protein autoprocessing reaction utilizing purified components. Instead, it is the response to *Shh* signaling that is affected, as indicated by failure of exogenously added Shh-N to induce ventral cell types in the presence of teratogenic compounds. Furthermore, although exogenously added Shh-N protein can induce endogenous *Shh* gene expression in neural plate explants (64, 65), we have demonstrated a complete inhibition of response by these teratogens at 2nM Shh-N, a concentration at which there is not induction of floor plate cells and therefore no endogenous *Shh* expression. The inhibitory effects of these compounds are dose-dependent, as demonstrated: (1) by maintenance or even expansion of the Isl-1 intermediate fate at intermediate inhibitor concentrations below those required for complete inhibition; and (2), by the requirement for correspondingly higher concentrations of teratogenic compounds to inhibit the response to increasing levels of Shh-N protein. A further indication of the specificity of these effects is the inability of these compounds to block cell behaviors such as migration, expression of Pax7, or HNK-1, or the response to other inductive signals such as BMP7 at concentrations that completely block the response to *Shh* signaling.

Our studies of sterol synthesis and transport suggest that these compounds are acting as class 2 inhibitors of cholesterol biosynthesis (61). For several reasons, however, simple reduction of cholesterol levels seem unlikely to account for the effects of these compounds on *Shh* signaling. First, the non-teratogenic compound tomatidine also displays potent inhibitory effects on cholesterol synthesis. Second *Shh* signaling in explants is not inhibited by 25-hydroxycholesterol, a hydroxysterol that blocks *de novo* cholesterol

biosynthesis (66). We can also rule out an inhibitory role for specific sterol precursors that may accumulate in drug-treated cells, since addition of 25-hydroxycholesterol together with inhibitory compounds should eliminate synthesis of sterol precursors yet does not restore the ability to respond to *Shh* signaling (67). An alternative mechanism to simple reduction of cholesterol would be a disruption of intracellular transport.

We have also shown that triparanol, jervine, and cyclopamine are potent inhibitors of PM cholesterol esterification, consistent with their classification as class 2 inhibitors. Consistent with transport disruption as the mechanism of drug action in inhibiting *Shh* signaling, we have found that several other previously characterized class 2 compounds also are able to inhibit the response to *Shh* signaling in explants (68). Tomatidine, however, also blocks esterification, indicating that general inhibition of this transport pathway is not sufficient for an inhibitory effect on the *Shh* response. We are currently investigating the possibility that this pathway comprises multiple steps that are differentially affected by tomatidine and the teratogenic compounds, and that only those steps not essential for the *Shh* response are affected by tomatidine. The unusual sterols that accumulate in tomatidine-treated cells are associated with peroxisomal sterol metabolism (60), consistent with such a differential effect of tomatidine on intracellular sterol transport.

In light of these drug effects on cholesterol homeostasis, it is interesting to note the presence of a sterol sensing domain (SSD) within Ptc, a key regulator of the *Shh* signaling pathway (33). The Ptc SSD initially was detected as a region of similarity to the Niemann-Pick C. Disease (NP-C) gene (31, 32). The similarity between Ptc and the NP-C protein extends beyond the five transmembrane spans of the SSD to include all twelve of the proposed transmembrane spans of Ptc. The significance of this sequence homology is not known, and the role of the SSD in NP-C is not clear, although this protein is proposed to regulate intracellular trafficking and loss of its function leads to lysosomal cholesterol accumulation (69). The SSDs of other proteins confer differential responses to high and low levels of intracellular sterols. The HMGCoA reductase enzyme thus displays a 3-5 fold decrease in stability as sterol concentration rise, and this behavior is dependent on the presence of the SSD. The SCAP regulator protein at low (but not at high) sterol concentrations stimulates the activity of the S2P metalloprotease, resulting in cleavage and activation of the SREBP transcription factor.

Those of the class 2 cholesterol synthesis inhibitors which have been examined appear to increase HMGCoA reductase activity and to stimulate the cleavage of SREBP. Given the localization of these two proteins to the ER, a likely mechanism for this effect is that disruption of sterol transport from PM to ER by class 2 compounds induces a 'low sterol' state in these ER proteins, despite higher levels of cellular sterols overall. The teratogenic compounds studied here all affect cholesterol synthesis and transport, and it is conceivable that they alter the normal distributions of sterols within intracellular compartments. If the function of Ptc is critically dependent upon the sterol concentrations in particular compartment, skewed sterol distributions in this compartment could act to perturb Ptc function via its SSD. One other possibility is that the function of Ptc in *Shh* signaling involves regulation of intracellular transport, as has been suggested for the related NP-C protein. If this were true, the perturbations of transport generated by these teratogenic compounds might affect the transport functions of Ptc in such a manner as to inhibit *Shh* signaling.

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40. Fertile chick eggs (white leghorn) were placed in a humidified incubator at 37.5° C in a rotating tray for 14 hours. The eggs were windowed at the air space and 250 µl of a sonicated 1 mg/ml jervine solution (Leibovitz's L15 medium, Gibco BRL) was applied under the shell membrane. The window was taped and the eggs incubated for an additional 4 days. The embryos were dissected in phosphate buffered saline (PBS, pH 7.2). The heads were removed from the trunk at the superior boarder of the heart and fixed in 3% Glutaraldehyde (EM grade, Polysciences, Inc.) in 0.1M sodium cacodylate (Polysciences, Inc.), 3 mM  $\text{CaCl}_2$  (pH 7.4) overnight at 4°C. They were washed in 0.1M sodium cacodylate (pH 7.4 placed in 2% osmium tetroxide (Polysciences, Inc.), 0.1M sodium cacodylate (pH 7.4) for 2 hours and washed in water. The samples were then dehydrated in a 50%, 70%, 90% and 100% ethanol series. Samples were critical point dried in liquid  $\text{CO}_2$  (CPD Model 10, Polaron), sputter coated with gold-palladium (Denton Desk II unit) and viewed on an Amray 1810 SEM operated at 20 kV.
41. Hamburger and Hamilton stage 9-10 (8-10 somites) embryos were used for all explant assays. Dissections were carried out in Leibovitz's L15 medium (Gibco BRL). Midline tissue just rostral to Hensen's node and well caudal to the last somite was removed with fine scissors. The neural ectoderm was separated from the lateral plate mesoderm and endoderm with dispase (Boehringer Mannheim, grade II 2.4 U/ml) treatment and then washed in L15. Midline, intermediate and ventral neural plate explants were further dissected with tungsten needles as diagrammed in figures 3A, 6A and 7A. Dissected tissues were transferred to a chambered coverglass (Nunc) in a drop of collagen (vitrogen 100, Collagen Biomaterials, Palo Alto, CA) containing 1X modified Eagle's medium (Gibco BRL) and 24 mM  $\text{NaH}_2\text{CO}_3$  (final pH 7.4-7.6), and warmed to 37.5°C for 30 minutes (in the absence of  $\text{CO}_2$ ) for gelation. Explants were cultured in 400 µl of F12 Nutrient Mixture (Ham) with glutamine (Gibco BRL), containing N-2 supplement (1X, Gibco BRL) and 100 U/ml penicillin and 100 ug/ml streptomycin in a 5%  $\text{CO}_2$ , humidified incubator at 37°C. AY 9944, triparanol, jervine, cyclopamine and tomatidine (all from 10mM stocks in 95% ethanol, except AY 9944 which is water soluble), purified Shh-N and BMP 7 were added at the initiation of the cultures. All of the explants were cultured for 40-48 hours except for the intermediate neural plate explants assayed for pax7 repression, which were cultured for 20-22 hours. At the end

- of the incubation period, explants were fixed in 4.0% formaldehyde (EM grade, Polysciences, Inc.) in PBS for 1 hour at 4°C, washed with PBS and then stained with a secondary antibody for 2 hours at room temperature. Rabbit anti-rat HNF3 $\beta$  (K2) 1:2000, mouse anti-ISL1 (40.2D6) 1:1000, mouse anti-pax7 1:10, mouse anti-rat HNK-1/N-CAM (sigma Biosciences) 1:1000, FITC-conjugated donkey anti-mouse IgG (Jackson ImmunoResearch Laboratories, Inc.) 1:100 and LRSC-conjugated donkey anti-rabbit IgG (Jackson ImmunoResearch Laboratories, Inc.) 1:300 were all diluted in PBTS. The explants were examined with an Olympus IX60 inverted microscope using a planapo objective with a 1.4 numerical aperture. Images were generated by confocal laser scanning microscopy with a cripton-argon laser exciting at 488 and 568 nm with emissions at 450-550 and 550-650 nm and utilizing Oz with Intervision software (Noran) on a Silicon Graphics Inc. platform.
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  47. HK293 cells, stably transfected with *Shh* using the Ecdysone-Inducible Mammalian Expression System (invitrogen), were plated in 6-well culture plates (Flacon, well area 9.6 cm<sup>2</sup>) in Dulbecco's modified Eagle's medium (DMEM, Gibco), 10% fetal bovine serum (FBS), 400  $\mu$ g/ml Zeocin Invitrogen), 2 mM L-glutamine, 100 U/ml Penecillin, 100  $\mu$ g/ml Stregtomycin, 350  $\mu$ g/ml G418 (Invitrogen) at 30-40% confluency and grown at 37°C. The following day, the media was changed to one that contained 10% dilapidated serum (K.M. Gibson et al, J. Lipid Res. **31**, 515 (1990)) and 1% ITS (Sigma) and otherwise was the same as above. After 24 hours, the cells were induced to express *Shh* with the addition of 1 $\mu$ M muristerone A (Invitrogen). AY 9944, triparanol, jervine, cyclopamine and tomatidine (all from 10mM stocks in 95% ethanol, except AY 9944 which is water soluble) were added to the cultures at the time of induction. The control cells received 0.475% ethanol to equal the maximum ethanol concentration in the 50  $\mu$ M steriodal alkaloid treatments. After an additional 24 hours,



- the culture supernatants were removed and the cells were lysed in the plate with 3X SDS-PAGE cell lysis buffer (3% SDS), diluted two-fold with water and boiled. Lysate samples (and in a separate experiment supernatant samples, for which the data is not shown) were loaded onto SDS-12% polyacrylamide gels for analysis, immunoblotted with primary antibodies for Shh-N and actin (Amersham) and horseradish peroxidase-conjugated secondary antibodies (Jackson ImmunoResearch Laboratories, Inc.), and visualized with a with luminescent substrate (Pierce).
48. Shh processing in transiently transfected cells is inefficient, with accumulation of 50-80% of Shh protein as unprocessed precursor. Even in these circumstances, we did not observe any effect of jervine, cyclopamine, or tomatidine upon Shh processing efficiency.
49. Unpublished data.
50. The in vitro studies of Hh autoprocessing used a bacterially expressed derivative of the *Drosophila* Hh protein (Porter 96A). The reactions were carried out as described (Porter 96B), except that the sterols and steroidal alkaloids were dried down from an ethanol or chloroform stock and resuspended in a 0.2% Triton-X 100 solution in a bath sonicator prior to addition to the reaction mixture.
51. Other sterols that participate in the reaction with similar efficiency to cholesterol are  $\beta$ -sitosterol, 5-androsten-3 $\beta$ -ol, ergosterol, 4 $\beta$ -hydroxycholesterol, 19-hydroxycholesterol, 20 $\alpha$ -hydroxycholesterol, 22(S)-hydroxycholesterol, 22(R)-hydroxycholesterol and 25-hydroxycholesterol. Epicholesterol, cholesterol acetate,  $\alpha$ -ecdysone, 20-OH ecdysone and thiocholesterol are unable to participate in the reaction.
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57. Pooled human lymphoblasts were washed with serum free RPMI-1640, then plated in 35 mm microwells in RPMI-1640 with 15% delipidated FBS (Gibson 90) and cultured at 37°C in a 5% CO<sub>2</sub> humidified atmosphere for 12 hours. AY 9944, triparanol, jervine, cyclopamine or tomatidine was then added and the cells were incubated for

- five days, after which the neutral sterols were extracted and analyzed as described by R.I. Kelley (Clin. Chim. Acta 236, 45 (1995)). Briefly, pelleted cells were saponified at 60°C in 4% (w/v) KOH in 90% ethanol with epicoprostanol as carrier, mixed with an equal volume of water and extracted three times in hexane. The hexane extracts were
- 5 dried under nitrogen, derivatized with bistrimethylsilyltrifluoroacetamide (BSTFA, Pierce) in pyridine and analyzed by selected ion monitoring gas chromatography/mass-spectrometry (SIM-GC/MS), utilizing a Hewlett Packard (HP) 5890A splitless injection port, a 0.2 mm x 25 m HP-1 methylsilicone (0.33 µm liquid phase) capillary column and a HP 5970A mass selective detector operated in electron impact mode at 70
- 10 eV with an ion source temperature of 200°C.
58. For determining their effects on sterol composition, AY 9944 and triparanol were used at 0.5µM and jervine, cyclopamine, and tomatidine were used at 10 µM. Doses lower than these produced more normal sterol profiles; higher doses increased the relative levels of cholesterol precursors but also reduced cell growth during the five day
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59. Sterols 1a, 1c-g, 2a,b and 3a,b are all intermediates in normal cholesterol biosynthesis, and 1b is thought to derive from 1a (G. Salen et al., J. Lipid Res. 37, 1169 (1996)).
60. Sterol 1h is associated with peroxisomal sterol synthesis and is particularly prominent in tomatidine treated cells. Sterol 4 is seen only in normal cells treated with tomatidine,
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- 25 63. Esterification of plasma membrane [<sup>3</sup>H] cholesterol in hepatoma cells was assayed according to Lange and Steck. Briefly, AH22 Hepatoma cells were cultured in 25 cm<sup>2</sup> flasks to ~89-90% confluency in DMEM 10% FBS at 37°C. The cells were washed in PBS and then labeled with 1.38 µCi [<sup>3</sup>H] cholesterol (3.17x10<sup>-5</sup>mmol cholesterol) in PBS for 10 minutes at 37°C. The [<sup>3</sup>H]cholesterol was in a vortexed solution of 2.5%
- 30 Triton WR-1339, 2.5 mM NaPi (pH 7.5) and 0.125 M sucrose. The cells were then washed in PBS with 0.5 mg/ml bovine serumalbumin (BSA) and incubated for 1.5 hours at 37°C in DMEM 10%FBS without or with AY 9944, triparanol, jervine,

cyclopamine or tomatidine. The cells were detached with trypsin, washed and suspended in 1 ml PBS. The sterols were then extracted with 2.5 ml of chloroform:methanol (2:1), dried on a speed vacuum concentrator, resuspended in 50  $\mu$ l of chloroform and spotted on silica gel G coated TLC plates (Merck). Cholesteryl esters and cholesterol were fractionated with a heptane:ether:acetic acid solvent (20:5:1), dried, visualized with I<sub>2</sub> vapor, scraped and counted directly in an aqueous scintillation counting cocktail (Econo-Safe, Research Products International Corp.)

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66. None of the explant responses to treatment with 2nM or 25 nM Shh-N were affected by additional addition of 25-OH cholesterol at 25  $\mu$ M. 25-OH cholesterol is a potent inhibitor of HMG CoA reductase and at the concentrations used blocks de novo cholesterol synthesis in chick embryos and in cultured cell systems (data not shown; S.C. Miller and G. Melnykovich, *J. Lipid Res.* **25**, 991 (1984); J.J. Bell, T.E. Sargeant and J.A. Watson, *J. Bio. Chem.* **251**, 1745 (1976)).

67. Addition of 25  $\mu$ M 25-hydroxycholesterol to explant cultures did not reverse the inhibitory effects of any of the teratogenic compounds.

68. Class 2 cholesterol synthesis inhibitors at the given concentrations block the response of intermediate neural plate explants to 25 nM Shh-N, without affecting signaling by BMP7:U 18666A 0.25  $\mu$ M, chloroquine 50  $\mu$ M, imipramine 75  $\mu$ M, progesterone 20  $\mu$ M.

69. P. G. Pentchev, et al., *Biochimica et Biophysica Acta* **1225**, 235-243 (1994).

#### Example 2: Essential role for Sonic *hedgehog* during hair follicle morphogenesis

The hair follicle is a source of epithelial stem cells and site of origin for several types of skin tumors. While it is clear that follicles arise by way of a series of inductive tissue interactions, identification of the signaling molecules driving this process remains a major challenge in skin biology. In this study we report an obligatory role for the secreted morphogen Sonic hedgehog (Shh) during hair follicle development. Hair germs, comprising epidermal placodes and associated dermal condensates were detected in both control and *Shh* <sup>-/-</sup> embryos, but progression through subsequent stages of follicle development was blocked in mutant skin. The expression of *Glil* and *Ptc1* was reduced in

*Shh*  $-/-$  dermal condensates and they failed to evolve into hair follicle papillae, suggesting that the adjacent mesenchyme is a critical target for placode-derived Shh. Despite the profound inhibition of hair follicle morphogenesis, late-stage follicle differentiation markers were detected in *Shh*  $-/-$  skin grafts, as well as cultured vibrissa explants treated with cyclopamine to block Shh signaling. Our findings reveal an essential role for Shh during hair follicle morphogenesis, where it is required for normal advancement beyond the hair germ stage of development.

## 10 Introduction

Early stages of organogenesis are marked by the appearance of mesenchymal condensates and focal cellular aggregates, or placodes, in adjacent epithelia. This process is driven to completion by a series of inductive signals traveling between epithelial and mesenchymal cell populations which ultimately give rise to the adult structure (reviewed in Gurdon, 1992; Thesleff *et al.*, 1995). In skin appendages such as vibrissae and hair follicles, detailed analysis of tissue recombinants has revealed the existence of at least three morphogenetic signals: the embryonic dermis instructs overlying ectoderm to initiate placode formation; the placode transmits a signal generating a dermal condensate with hair follicle-inductive properties; and the condensate in turn sends a signal to nascent follicle keratinocytes stimulating their proliferation, downgrowth into the developing dermis, and reorganization to form the mature follicle (reviewed in Sengel, 1976; Hardy, 1992). The epithelial and mesenchymal components of the follicle remain in close proximity in mature hair bulbs, where the dermal papilla is surrounded by matrix cells giving rise to at least six phenotypically distinct epithelial cell types in the hair shaft and inner root sheath of the hair follicle. After birth the follicle epithelium cycles through periods of active growth (anagen), followed by regression (catagen) and inactivity (telogen) (reviewed in Cotsarelis, 1997). The morphogenetic program that accompanies the transition from telogen to anagen bears similarities to follicle development during embryogenesis, making this structure a unique model for studying certain aspects of organogenesis in the adult animal. Although a large number of genes have been implicated at various stages of hair follicle development and cycling (reviewed in Rosenquist and Martin, 1996; Stern *et al.*, 1996; Widelitz *et al.*,

1997; Millar, 1997), the molecular nature of the inductive signals that underlie the formation of the follicle is largely unknown.

*In situ* localization of transcripts encoding potential morphogens has revealed focal expression of *Sonic hedgehog* (*Shh*) in placodes of the epidermis and several other epithelia at early stages of development, with *Ptcl* transcripts encoding a putative Shh receptor also present in adjacent mesenchymal cells (Bitgood and McMahon, 1995; Iseki *et al.*, 1996; Oro *et al.*, 1997; Motoyama *et al.*, 1998). These findings, coupled with the accumulating evidence demonstrating a pivotal role for secreted Hedgehog proteins in a variety of developmental processes (reviewed in Hammerschmidt *et al.*, 1997), led us to examine the potential involvement of this pathway in hair follicle morphogenesis. Since the follicle is a source of cutaneous stem cells and a likely site of origin for certain epithelial skin cancers (Cotsarelis *et al.*, 1990; Lavker *et al.*, 1993; Rochat *et al.*, 1994; Hansen and Tennant, 1994), understanding the developmental biology of this organ is likely to provide insights relevant to normal skin function as well as wound-healing and neoplasia, and may shed light on fundamental aspects of organogenesis involving other structures as well.

## Methods

### Animals and Skin Transplantation

The generation and identification of *Shh* mutant mice was performed as described (Chiang *et al.*, 1996). Embryonic skin was grafted onto the dorsal fascia of nude mice beneath a protective silicone chamber using a modification of a previously-described technique (Dlugosz *et al.*, 1995). The chamber was removed 11-12 days after grafting and tissue harvested for analysis after an additional one to four weeks. Animals were handled in accordance with NIH guidelines.

### Immunohistochemistry

Tissue was fixed overnight in Carnoy's or Bouin's solution for detecting keratins, (KI, K10, K5, K14, and K 17), loricrin, and filaggrin; fixation with neutral-buffered formalin was used for tissues immunostained with Lef-1, Ki67, and hair keratin (AE13)

antibodies. Samples were embedded in paraffin and 8  $\mu$ m sections cut for immunostaining. Immunoreactivity of antigens in formalin-fixed sections was restored by immersing slides in a boiling 0.01M citrate buffer, pH 6, for 10 minutes. The following primary antibodies were used at the indicated dilutions for immunostaining: rabbit anti-keratins K 1, K 10, K5 and K 14 (1:500) (Roop *et al.*, 1984), loricrin and filaggrin (1:500) (Roop *et al.*, 1987), supplied by Dr. Stuart Yuspa; rabbit anti-K17 (1:1000) (McGowan and Coulombe, 1998), provided by Dr. Pierre Coulombe; rabbit anti-Lef-1 (1:200) (Travis *et al.*, 1991), a gift from Dr. Rudolf Grosschedl; rabbit anti-Ki67, NCL-Ki67p (Novocastra Laboratories, Ltd., Newcastle upon Tyne, UK) (1:200); and mouse monoclonal AE 13 hybridoma supernatant, which recognize type I hair keratins (1:5) (Lynch *et al.*, 1986), provided by Dr. Tung-Tien Sun. Tissue sections were incubated with primary antibodies diluted in tris-buffered saline containing 1% bovine serum albumin, typically for 1-2 hours at room temperature. Subsequent immunostaining procedures were performed using peroxidase Vectastain ABC kits (Vector Laboratories, Inc., Burlingame, CA) and 3,3'-diaminobenzidine (Sigma, St. Louis, MO) as a substrate, according to the manufacturers' recommendations. Sections were counterstained with hematoxylin and mounted using Permount (Fisher Scientific, Pittsburgh, PA).

#### In Situ Hybridization

Non-radioactive RNA *in situ* hybridization was performed on 5  $\mu$ m sections essentially as described (Groves *et al.*, 1995), using previously described sequences for *Glil* (Walterhouse *et al.*, 1993), *Ptcl* (Goodrich *et al.*, 1996), and *BMP-4* (Jones *et al.*, 1991).

#### Vibrissa Follicle Explants

Vibrissa follicle explants were established using CD-1 mouse embryos at 13.5 days of gestation according to a previously described protocol (Hirai *et al.*, 1989), with minor modifications. Vibrissa pads were transferred onto Nuclepore filters (13 mm, 8  $\mu$ m pores), and floated on, 2 ml of medium [DMEM (Life Technologies, Gaithersburg, MD) + Ham's F12 medium (Life Technologies) (1:1), with 1% FCS (Intergen, Purchase, NY), penicillin (50 units/ml) and streptomycin (50  $\mu$ g/ml) (Life Technologies)] in 6-well plates. Similar

results were obtained using a DMEM-based medium, without the addition of Ham's F12. Explants were fed fresh medium every two days. Microdissection was performed with the aid of a Nikon SMZ-2T stereomicroscope and photomicrographs were taken using an Olympus OM-4 camera. Cyclopamine was stored at -20 as a 10 mM stock in 95% EtOH.

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#### RNA Isolation and RT-PCR

RNA was obtained by solubilizing individual explants in TriZol (Life Sciences) and isolating as recommended by the manufacturer. cDNA was synthesized using SuperScript II Rnase H reverse transcriptase with random primers (Life Technologies), and RT-PCR performed using the following primers: MHKA1 (318 bp product), (forward 5'-ATCAGAGAATGCCAGGTTGG-3' and reverse 5'-TCATTGAGCACACGGTTCAG-3'); hacl-1 (308 bp product), (forward 5'-TTGTATCTCCACTCCTGCCC.3 and reverse 5'-AGACTCCACAGGTTGGTTGG-3'); profilaggrin (330 bp product), (forward 5'-GCTTAAATGCATCTCCAG-3' and reverse 5'-AGTCAGTCCTATTGCAGG-3') (Bickenbach *et al.*, 1995); P actin (421 bp product), (for-ward 5'-TACCACAGGCATTGTGATGGA-3' and reverse 5'-CAACGTCACACTTCATGATGG-3') (Walterhouse *et al.*, 1993). The following PCR conditions were used for MHKA1, Hacl-1, and actin: 95 x 3 min "hot start"; 95 x 50 sec, 58 x 30 sec, and 72 x 60 sec for 25 (actin) or 35 cycles (MHKA I and Hacl-1); 72 x 7 min. PCR conditions for profilaggrin primers were as previously described (Bickenbach *et al.*, 1995). Reaction products were run through 1.5% agarose gels and visualized with ethidium bromide.

#### **Results and Discussion**

Early stages of hair follicle development appeared similar in control and *Shh*<sup>-/-</sup> embryos. Hair germs, consisting of clusters of columnar basal keratinocytes protruding into the developing dermis with associated dermal condensates, were detected in the skin of both mutant and control embryos at 15.5 days of gestation (Fig. 8 A,B). Despite the similar morphology of control and *Shh*-deficient hair germs, a dramatic difference in gene expression patterns was revealed by *in situ* hybridization. The level of *Glil* mRNA was markedly reduced in both the epithelial and mesenchymal components of *Shh*<sup>-/-</sup> primary hair germs (Fig. 8C,D). In addition, expression of *Ptcl* was reduced in *Shh* mutant hair

germs, although some placodes contained levels slightly above background (Fig. 8E,F). These findings are consistent with previous reports identifying Shh as a positive regulator of both *Glil* and *Ptcl* (Marigo and Tabin, 1996; Marigo *et al.*, 1996; Lee *et al.*, 1997; Sasaki *et al.*, 1997), and suggest that Shh is signaling in both the epithelial and mesenchymal cells of the developing follicle. In contrast to *Glil* and *Ptcl*, *BMP-4* mRNA was clearly detectable in condensates of mutant and control embryos (Fig. 8G,H), arguing against a requirement for Shh in the induction of *BMP-4* expression. Thus, although Shh is not required for the initiation of hair follicle development, primary hair germs that arise in Shh mutant skin are deficient in the expression of at least some Shh target genes.

In control embryos, the interval between E15.5 and E17.5 is marked by rapid proliferation and downgrowth of the follicle into the developing dermis, accompanied by a several-fold increase in the mass of the follicle epithelium and reorganization into distinct cellular compartments. In the most mature follicles, keratinocytes in the most peripheral cell layer, which give rise to the outer root sheath in the mature follicle, have assumed a columnar arrangement perpendicular to the long axis of the developing follicle; cells located centrally are without a definite orientation at this stage but will eventually be replaced by the three concentric layers of inner root sheath cells and the three cell types comprising the hair shaft; and the epithelial cells of the deepest portion of the follicle, the future hair bulb, have surrounded what is at this stage a well-defined cluster of mesenchymal cells, the dermal papilla (Fig. 9A, arrow). Even the less mature follicles exhibit an organized "cap" of mesenchymal cells at their invaginating tips (Fig. 9A, arrowheads). In striking contrast, hair follicles in skin from mutant embryos at E 17.5 failed to develop past the hair germ stage seen at E 15.5 (Fig. 9B). Although the follicle epithelium was most obviously affected due to its lack of growth, organizing dermal condensates and dermal papillae were conspicuously absent in mutant skin. These results are consistent With the idea that epidermis-derived Shh (Bitgood and McMahon, 1995; Iseki *et al.*, 1996; Oro *et al.*, 1997; Motoyama *et al.*, 1998) functions as a paracrine signal regulating development of the mesenchymal component of the hair follicle. Inhibition of follicle formation is not likely to be due to a general disruption of skin development since epidermal morphogenesis, marked by the appearance of granular and cornified cell layers, took place by E 17.5 in both control and mutant embryos (Fig. 9A,B).



Additional studies were performed to determine whether Shh influenced the expression of epithelial differentiation markers in embryonic skin. Keratinocytes in developing hair follicles can be distinguished by a relative deficiency of K5 and K14, keratins that are abundant in surrounding epidermal basal cells (Kopan and Fuchs, 1989; Byrne *et al.*, 1994). Immunohistochemical staining of E17.5 embryos revealed greatly reduced or undetectable levels of K14 in a sub-population of cells comprising the normal follicles in control embryos as well as the primordial follicles seen in *Shh*<sup>-/-</sup> embryos (Fig. 9C,D; arrows). Moreover, K17, which is normally not detected in interfollicular epidermis but is expressed in developing and mature hair follicles (Panteleyev *et al.*, 1997; McGowan and Coulombe, 1998), was localized to the follicular epithelium in both control and mutant skin (Fig. 9E,F). Thus, although morphogenesis of hair follicles in *Shh*<sup>-/-</sup> skin fails to progress past the hair germ stage, these structures contain epithelial cells that have initiated a terminal differentiation program characteristic of developing follicle keratinocytes. Consistent with the morphological findings in Figs. 9A and B, the expression level of epidermal-specific differentiation markers (keratins 1 and 10, loricrin, and filaggrin) in *Shh*<sup>-/-</sup> skin was similar to or greater than in control epidermis, based on immunohistochemical staining (data not shown).

Since *Shh*<sup>-/-</sup> mice are not viable, post-natal analysis of mutant skin was performed following grafting onto nude mice. Whereas skin from control mice produced abundant pigmented hairs, transplanted *Shh*<sup>-/-</sup> skin failed to generate detectable hairs but exhibited a pigmented graft site, consistent with the strain of donor skin (Fig. 10A). The histology of control skin grafts revealed the typical structures seen in normal mouse skin, including numerous hair follicles and sebaceous glands (Fig. 10B). In striking contrast, mutant skin failed to produce normal-appearing follicles, hair shafts, or sebaceous glands, but in some cases (3 of a total of 7 *Shh*<sup>-/-</sup> grafts), exhibited a thickened epidermis with focal areas of hyperkeratosis (Fig. 10C). Conspicuous aggregates of basophilic cells with scant cytoplasm were detected at the dermal-epidermal junction in these mutant grafts (Fig. 10C, arrows). Interestingly, the morphology of cells in the Shh-deficient keratinocyte aggregates was reminiscent of cells in control hair bulbs, and additional analyses revealed biochemical similarities. Cells in these aggregates were unreactive with K5 antibodies (Fig. 10D, arrows), exhibited abundant nuclear Lef-1 expression (Fig. 10 E) (Zhou *et al.*, 1995), and contained a high proportion of proliferating cells detected by Ki67

immunostaining (data not shown). Interestingly, short columnar structures resembling abortive hair shafts were associated with some of the *Shh* mutant keratinocyte aggregates. Moreover, these structures expressed hair-specific keratin (Fig. 10F), indicating that an advanced stage in the follicle differentiation program was achieved despite a dramatic  
 5 disruption of normal morphogenesis. Rarely, a small cluster of mesenchymal cells was seen associated with the base of a keratinocyte aggregate, as illustrated in Fig. 10E, where these cells immunostain with Lef-1 antibody. These findings suggest that a rudimentary dermal papilla is present in at least some of the hair germs seen in *Shh* mutant grafts.

To better define the temporal requirement for Shh during follicle development,  
 10 tissue culture studies were performed using cyclopamine (GaTfield and Keeler, 1996), which has recently been shown to block Shh signaling in neural plate explants (Cooper *et al.*, 1998; Incardona *et al.*, 1998). Explants were established using vibrissa pads obtained from mice at 13.5 days of gestation (Hirai *et al.*, 1989). When grown for six to eight days in culture, explants undergo robust morphogenesis resulting in the formation of elongated,  
 15 grossly normal-appearing vibrissa follicles (Fig. 11A). These follicles contained hair shafts and expressed genes encoding mouse hair keratin A I (MHKA 1) (Kaytes *et al.*, 1991) and a hair cortex-specific marker Hactl-1 (Huh *et al.*, 1994), detected by RT-PCR (Fig. 11B). Treatment of explants with cyclopamine results in striking inhibition of morphogenesis, indicating that Shh signaling is required during or shortly after the hair germ stage of  
 20 vibrissa follicle development (Fig. 11C). In keeping with our results obtained using *Shh* mutant skin, hair-specific transcripts are detected in cyclopamine-treated grafts (Fig. 11B) despite their altered development, providing further support for the notion that biochemical differentiation of the follicle is not necessarily coupled to its morphogenesis. Both control and cyclopamine-treated explants accumulate profilaggrin mRNA, indicating  
 25 that disruption of Shh signaling does not inhibit epidermal differentiation.

Collectively, the results of our studies reveal an obligatory role for Shh in the progression of hair follicle morphogenesis past the hair germ stage of development. The reduced expression of *Ptc1* and *Glil* in *Shh*  $-/-$  dermal condensates, coupled with their failure to evolve into recognizable dermal papillae, argue that Shh is involved in regulating  
 30 development of the mesenchymal component of the hair follicle, although a requirement for Shh signaling in the epithelial component of the follicle cannot be excluded. In the absence of dermal papillae normal hair follicle morphogenesis does not proceed,

underscoring the critical influence these cells have on growth and remodeling of developing follicle epithelium (Jahoda *et al.*, 1984; Weinberg *et al.*, 1993). Interestingly, biochemical differentiation of the follicle can take place in the absence of normal morphogenesis, implying that these two processes are regulated independently in this organ. Additional experiments will be required to formally define which component of the developing follicle is functionally impaired in *Shh*<sup>-/-</sup> embryos, and to determine whether Shh has additional roles at later stages of follicle development or during hair cycling. We anticipate that these studies may ultimately help explain how constitutive activation of the Shh signaling pathway in keratinocytes contributes to the formation of basal cell carcinoma (Johnson *et al.*, 1996; Hahn *et al.*, 1996; Oro *et al.*, 1997; Fan *et al.*, 1997; Xie *et al.*, 1998).

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### Example 3: Rescue of *ptc* loss-of-function phenotype

Based on the results presented above, we have attempt to determine the site in the *Shh* signaling pathway at which cyclopamine operates, and therefor better understand the spectrum of tumors caused by *Shh* pathway-activating lesions that could potentially be treated with this compound.

These studies involve the use of mouse embryonic fibroblasts (MEFs) that were generated by trypsin digestion of E8.5 embryos from patched (*ptc*) +/- matings. The mouse *ptc* gene was disrupted by homologous recombination in which part of exon1 and all of exon 2 were replaced with the bacterial *lacZ* gene (Goodrich *et al*, (1997) *Science* 277:1109). As *Ptc* protein suppresses *Shh* signaling, a loss of its function activates the *Shh* signaling pathway. *Shh* signaling, through a cascade of events, is mediated by the Gli transcription factors. One of the target genes of *Shh* signaling is *ptc*, through Gli-binding sites in the *ptc* promoter region, and this serves as a feedback mechanism for down regulation of signaling. Thus, in these *ptc* -/- embryos, the *Shh* signaling pathway is activated in many tissues, and the *lacZ* gene product  $\beta$ -galactosidase is expressed in all of those tissues as a report of pathway activation.

We obtained these MEFs to determine whether cyclopamine acts on *Ptc* or another, component of the cascade to inhibit *Shh* signaling. If the target of cyclopamine is *Ptc*, then one would expect that when the *Shh* pathway is activated by the loss of *ptc* function, it



could no longer be inhibited by cyclopamine. Figure 12 demonstrates that the Shh signaling pathway can be activated in these fibroblasts in cell culture, and that the level of  $\beta$ -galactosidase activity does reflect the degree of pathway activation. The MEF line 23-4 is heterozygous for *ptc-lacZ*, and thus contains one functional *ptc* allele capable of maintaining a repressed state of the pathway, but will express *lacZ* when the pathway is activated by addition of Shh protein (see Figure 12).

In contrast, the  $\beta$ -galactosidase activity in MEFs homozygous for *ptc-lacZ*, (cell line 23-1) is markedly elevated, because in these cells the pathway is constitutively activated by the loss of a functional *ptc* allele (Figure 13). When these cells are cultured with cyclopamine,  $\beta$ -galactosidase activity is decreased, indicating that when the Shh signaling pathway is unregulated by Ptc repression, it is still sensitive to cyclopamine inhibition. The reduction of  $\beta$ -galactosidase activity appears to result from the specific inhibition of Shh signaling, rather than from cell toxicity because enzymatic activity is normalized to whole protein content of the sample. Also, the reduction of  $\beta$ -galactosidase activity can be obtained with exposure to cyclopamine over a period of time that is shorter than the average cell cycle, and so does not appear to be due solely to an inhibition of cell proliferation (Figure 14).

A final indication that this represents specific inhibition of Shh signaling is that it cannot be achieved with a non-inhibitory, but structurally related compound tomatidine (Figure 15).

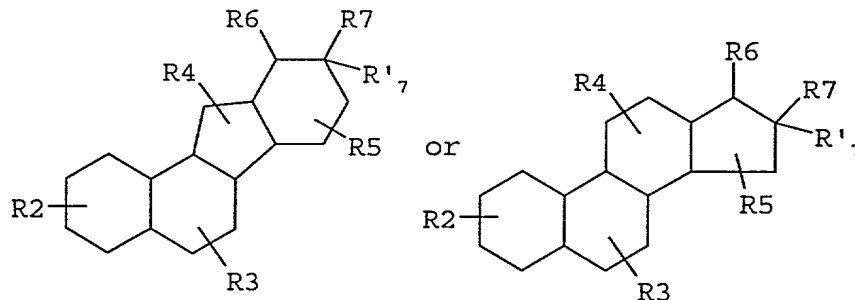
All of the references cited above are hereby incorporated by reference herein.

### *Equivalents*

Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the following claims.

## Claims

1. A method for inhibiting paracrine and/or autocrine signals produced by a hedgehog proteins comprising contacting a cell sensitive to the hedgehog protein with a  
5 hedgehog antagonist in a sufficient amount to reduce the sensitivity of the cell to the hedgehog protein, wherein the hedgehog antagonist is a organic molecule having a molecule weight less than 750 amu.
2. A method for inhibiting an altered growth state of a cell having a *ptc* loss-of-function phenotype or a smoothened gain-of-function phenotype, comprising  
10 contacting the cell with a *ptc* agonist in a sufficient amount to inhibit the altered growth state, wherein the *ptc* agonist is a organic molecule having a molecule weight less than 750 amu.
3. The method of claim 1 or 2, wherein hedgehog antagonist is a the steroidal alkaloid represented in the general formulas (I), or unsaturated forms thereof and/or seco-,  
15 nor- or homo-derivatives thereof:

Formula I

wherein, as valence and stability permit,

- 20  $R_2$ ,  $R_3$ ,  $R_4$ , and  $R_5$ , represent one or more substitutions to the ring to which each is attached, for each occurrence, independently represent hydrogen, halogens, alkyls, alkenyls, alkynyls, aryls, hydroxyl, =O, =S, alkoxyl, silyloxy, amino, nitro, thiol, amines, imines, amides, phosphoryls, phosphonates, phosphines, carbonyls, carboxyls, carboxamides, anhydrides, silyls, ethers, thioethers, alkylsulfonyls, arylsulfonyls, selenoethers, ketones, aldehydes, esters, or  $-(CH_2)_m-R_8$ ;

$R_6$ ,  $R_7$ , and  $R'_7$ , are absent or represent, independently, halogens, alkyls, alkenyls, alkynyls, aryls, hydroxyl, =O, =S, alkoxyl, silyloxy, amino, nitro, thiol, amines, imines, amides, phosphoryls, phosphonates, phosphines, carbonyls, carboxyls, carboxamides, anhydrides, silyls, ethers, thioethers, alkylsulfonyls, arylsulfonyls, selenoethers, ketones, aldehydes, esters, or  $-(CH_2)_m-R_8$ , or

$R_6$  and  $R_7$ , or  $R_7$  and  $R'_7$ , taken together form a ring or polycyclic ring, e.g., which is substituted or unsubstituted,

with the proviso that at least one of  $R_6$ ,  $R_7$ , or  $R'_7$  is present and includes a primary or secondary amine;

$R_8$  represents an aryl, a cycloalkyl, a cycloalkenyl, a heterocycle, or a polycycle; and

$m$  is an integer in the range 0 to 8 inclusive.

4. The method of claim 3, wherein:

$R_2$  and  $R_3$ , for each occurrence, is an -OH, alkyl, -O-alkyl, -C(O)-alkyl, or -C(O)- $R_8$ ;

$R_4$ , for each occurrence, is an absent, or represents -OH, =O, alkyl, -O-alkyl, -C(O)-alkyl, or -C(O)- $R_8$ ;

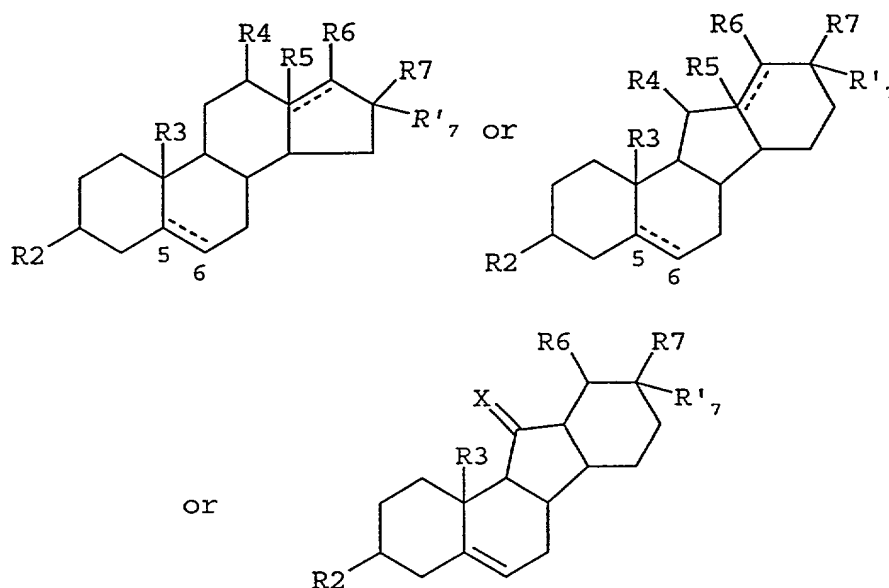
$R_6$ ,  $R_7$ , and  $R'_7$  each independently represent, hydrogen, alkyls, alkenyls, alkynyls, amines, imines, amides, carbonyls, carboxyls, carboxamides, ethers, thioethers, esters, or  $-(CH_2)_m-R_8$ , or

$R_7$ , and  $R'_7$  taken together form a furanopiperidine, such as perhydrofuro[3,2-b]pyridine, a pyranopiperidine, a quinoline, an indole, a pyranopyrrole, a naphthyridine, a thiofuranopiperidine, or a thiopyranopiperidine

with the proviso that at least one of  $R_6$ ,  $R_7$ , or  $R'_7$  is present and includes a primary or secondary amine;

$R_8$  represents an aryl, a cycloalkyl, a cycloalkenyl, a heterocycle, or a polycycle, and preferably  $R_8$  is a piperidine, pyrimidine, morpholine, thiomorpholine, pyridazine,

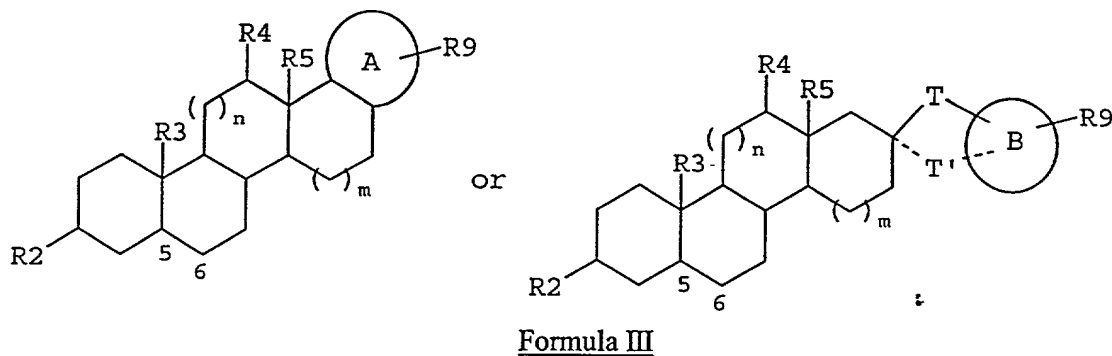
- 5 5. The method of claim 1 or 2, wherein the hedgehog antagonist is a steroidal alkaloid represented in the general formula (II), or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



10 Formula II

wherein  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ ,  $R_6$ ,  $R_7$ , and  $R'_7$  are as defined above, and X represents O or S, though preferably O.

- 15 6. The method of claim 1 or 2, wherein the steroidal hedgehog antagonist is an alkaloid represented in the general formula (III), or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



wherein

$R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$  and  $R_8$  are as defined above;

A and B represent monocyclic or polycyclic groups;

T represent an alkyl, an aminoalkyl, a carboxyl, an ester, an amide, ether or amine linkage of 1-10 bond lengths;

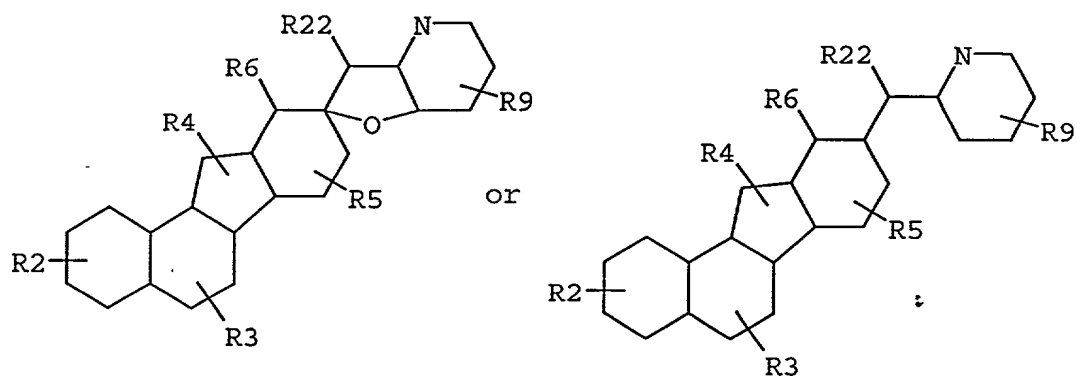
T' is absent, or represents an alkyl, an aminoalkyl, a carboxyl, an ester, an amide, ether or amine linkage of 1-3 bond lengths, wherein if T and T' are present together, than T and T' taken together with the ring A or B form a covalently closed ring of 5-8 ring atoms;

$R_9$  represent one or more substitutions to the ring A or B, which for each occurrence, independently represent halogens, alkyls, alkenyls, alkynyls, aryls, hydroxyl, =O, =S, alkoxyl, silyloxy, amino, nitro, thiol, amines, imines, amides, phosphoryls, phosphonates, phosphines, carbonyls, carboxyls, carboxamides, anhydrides, silyls, ethers, thioethers, alkylsulfonyls, arylsulfonyls, selenoethers, ketones, aldehydes, esters, or  $-(CH_2)_m-R_8$ ; and

n and m are, independently, zero, 1 or 2;

with the proviso that A and  $R_9$ , or T, T' B and  $R_9$ , taken together include at least one primary or secondary amine.

7. The method of claim 1 or 2, wherein the hedgehog antagonist is a steroidal alkaloid represented in the general formula (IV), or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:

Formula IV

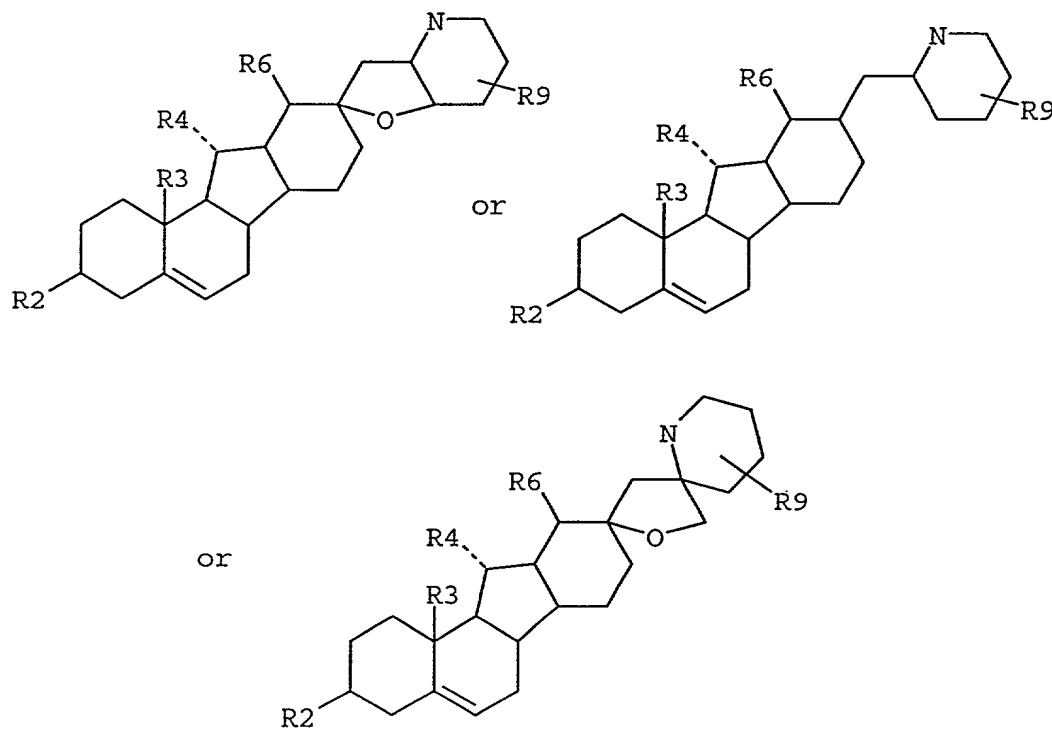
wherein

R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>5</sub>, R<sub>6</sub> and R<sub>9</sub> are as defined above;

5

R<sub>22</sub> is absent or represents an alkyl, an alkoxyl or -OH.

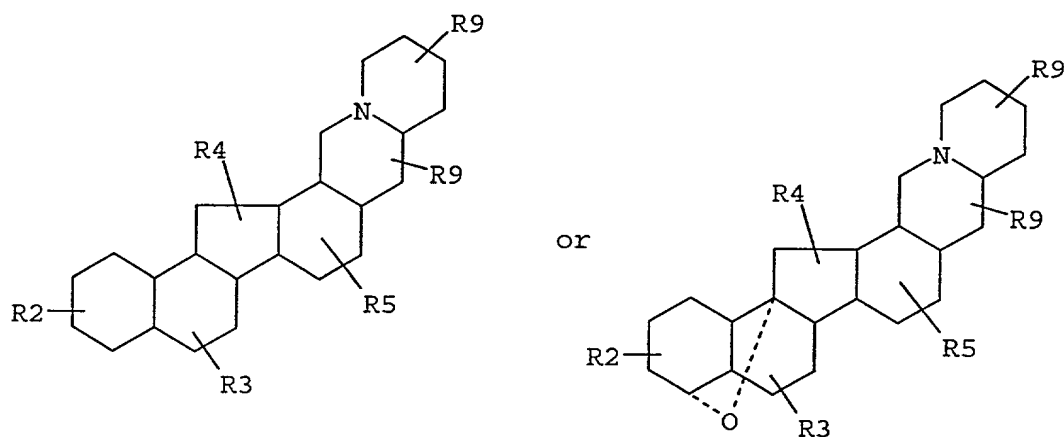
8. The method of claim 1 or 2, wherein the hedgehog antagonist is a steroidal alkaloid represented in the general formula (V) or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:

Formula V

10

wherein  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_6$  and  $R_9$  are as defined above;

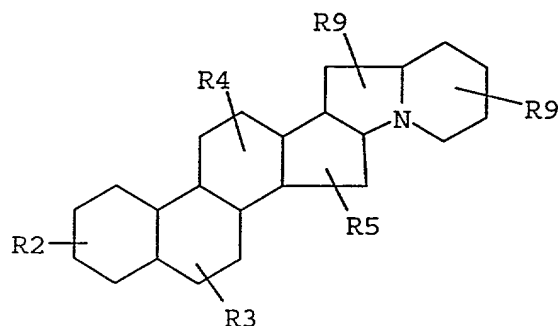
9. The method of claim 1 or 2, wherein the hedgehog antagonist is a steroidal alkaloid represented in the general formula (VI), or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula VI

wherein  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$  and  $R_9$  are as defined above;

10. The method of claim 1 or 2, wherein the hedgehog antagonist is a steroidal alkaloid represented in the general formula (VII) or unsaturated forms thereof and/or seco-, nor- or homo-derivatives thereof:



Formula VII

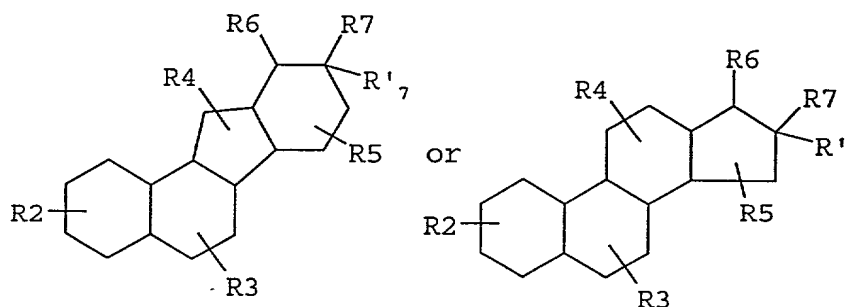
wherein  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$  and  $R_9$  are as defined above.

11. The method of any of claims 3-10, wherein the steroidal alkaloid does not substantially interfere with the biological activity of such steroids as aldosterone, androstane, androstene, androstenedione, androsterone, cholecalciferol, cholestane, cholic acid, corticosterone, cortisol, cortisol acetate, cortisone, cortisone acetate, deoxycorticosterone, digitoxigenin, ergocalciferol, ergosterol, estradiol-17- $\alpha$ , estradiol-17- $\beta$ , estriol, estrane, estrone, hydrocortisone, lanosterol, lithocholic acid, mestranol,  $\beta$ -methasone, prednisone, pregnane, pregnenolone, progesterone, spironolactone, testosterone, triamcinolone and their derivatives.
12. The method of any of claims 3-10, wherein the steroidal alkaloid does not specifically bind a nuclear hormone receptor.
13. The method of any of claims 3-10, wherein the steroidal alkaloid does not specifically bind estrogen or testosterone receptors.
14. The method of any of claims 3-10, wherein the steroidal alkaloid has no estrogenic activity at therapeutic concentrations.
15. The method of any of claims 1-10, wherein the hedgehog antagonist inhibits hedgehog-mediated signal transduction with an ED<sub>50</sub> of 1mM or less.
16. The method of any of claims 1-10, wherein the hedgehog antagonist inhibits hedgehog-mediated signal transduction with an ED<sub>50</sub> of 1 $\mu$ M or less.
17. The method of any of claims 1-10, wherein the steroidal alkaloid inhibit hedgehog-mediated signal transduction with an ED<sub>50</sub> of 1nM or less.
18. The method of claims 1 or 2, wherein the cell is contacted with the hedgehog antagonist *in vitro*.
19. The method of claims 1 or 2, wherein the cell is contacted with the hedgehog antagonist *in vivo*.
20. The method of claim 1 or 2, wherein the hedgehog antagonist is administered as part of a therapeutic or cosmetic application.
21. The method of claim 19 or 20, wherein the hedgehog antagonist is administered to treat a condition selected from the group consisting of regulation of neural tissues,



bone and cartilage formation and repair, regulation of spermatogenesis, regulation of smooth muscle, regulation of lung, liver and other organs arising from the primitive gut, regulation of hematopoietic function, regulation of skin and hair growth, etc.

- 5 22. The method of any of claims 1-7, wherein the hedgehog antagonist is applied as a topical formulation to skin in order to inhibit aberrant proliferation of epithelial cells.
23. The method of any of claims 1-17 or 22, wherein the hedgehog antagonist is administered to patient to inhibit growth of a basal cell carcinoma.
- 10 24. A pharmaceutical preparation comprising steroidal alkaloid is represented in the general formulas (I), or unsaturated forms thereof and/or seco-, nor- or homo- derivatives thereof:



Formula I

- 15 wherein, as valence and stability permit,

20  $R_2$ ,  $R_3$ ,  $R_4$ , and  $R_5$ , represent one or more substitutions to the ring to which each is attached, for each occurrence, independently represent hydrogen, halogens, alkyls, alkenyls, alkynyls, aryls, hydroxyl, =O, =S, alkoxyl, silyloxy, amino, nitro, thiol, amines, imines, amides, phosphoryls, phosphonates, phosphines, carbonyls, carboxyls, carboxamides, anhydrides, silyls, ethers, thioethers, alkylsulfonyls, arylsulfonyls, selenoethers, ketones, aldehydes, esters, or  $-(CH_2)_m-R_8$ ;

$R_6$ ,  $R_7$ , and  $R'_7$ , are absent or represent, independently, halogens, alkyls, alkenyls, alkynyls, aryls, hydroxyl, =O, =S, alkoxyl, silyloxy, amino, nitro, thiol,

amines, imines, amides, phosphoryls, phosphonates, phosphines, carbonyls, carboxyls, carboxamides, anhydrides, silyls, ethers, thioethers, alkylsulfonyls, arylsulfonyls, selenoethers, ketones, aldehydes, esters, or  $-(CH_2)_m-R_8$ , or

$R_6$  and  $R_7$ , or  $R_7$  and  $R'_7$ , taken together form a ring or polycyclic ring, e.g., which is substituted or unsubstituted,

with the proviso that at least one of  $R_6$ ,  $R_7$ , or  $R'_7$  is present and includes a primary or secondary amine;

$R_8$  represents an aryl, a cycloalkyl, a cycloalkenyl, a heterocycle, or a polycycle; and

$m$  is an integer in the range 0 to 8 inclusive.

25. The preparation of claim 24, formulated for topical application.

26. A process for manufacturing a medicament comprising formulating a steroid alkaloid inhibitor of a hedgehog signal transduction pathway in a pharmaceutically acceptable excipient to form a sterile medicament for preventing growth of cells having an aberrant activation hedgehog pathway.

**Abstract**

The present invention makes available assays and reagents inhibiting paracrine and/or autocrine signals produced by a hedgehog protein or aberrant activation of a hedgehog signal transduction pathway, e.g., which involve the use of a steroidal alkaloid or other small molecule.

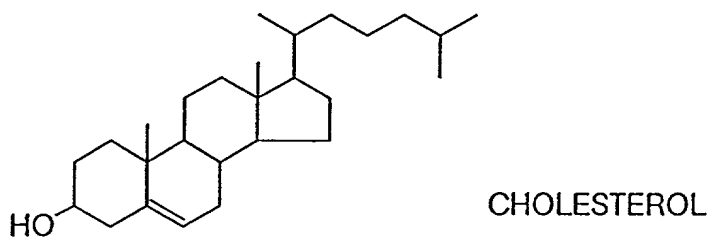
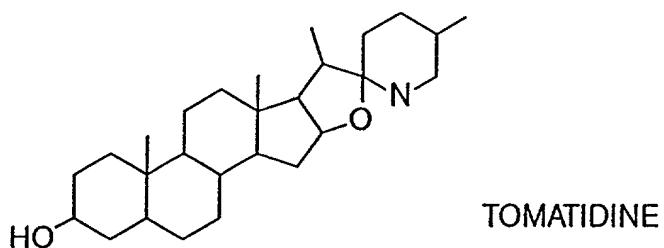
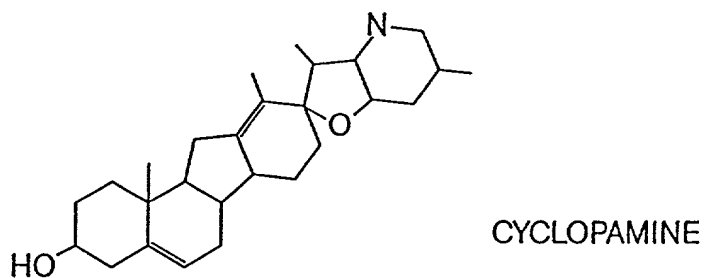
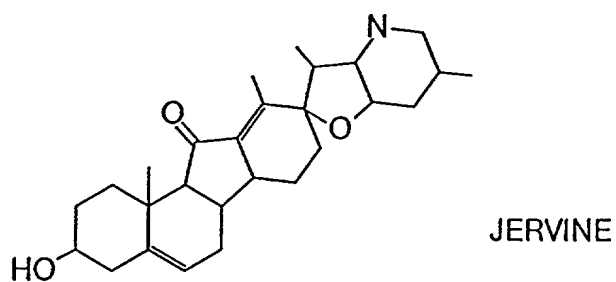
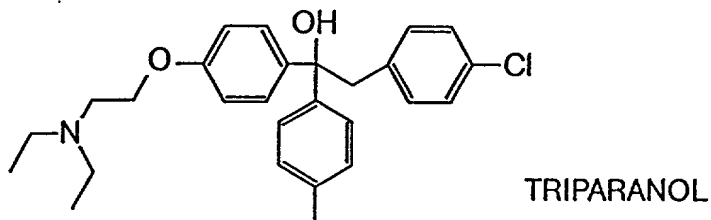
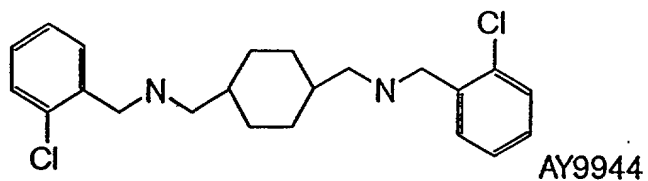


Fig. 1



Fig. 2A

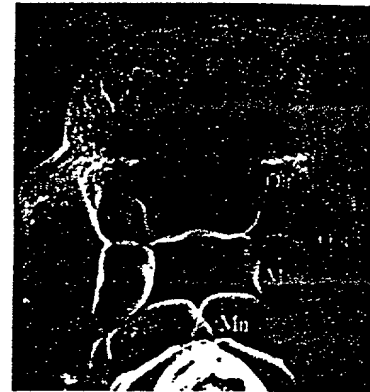


Fig. 2B



Fig. 2C



Fig. 2D

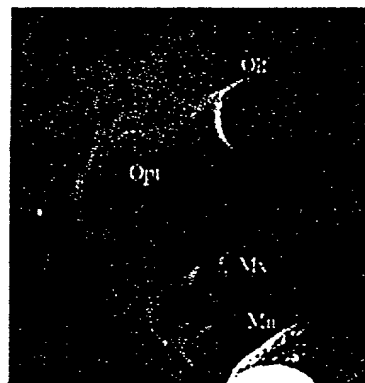


Fig. 2E

008077-12630260

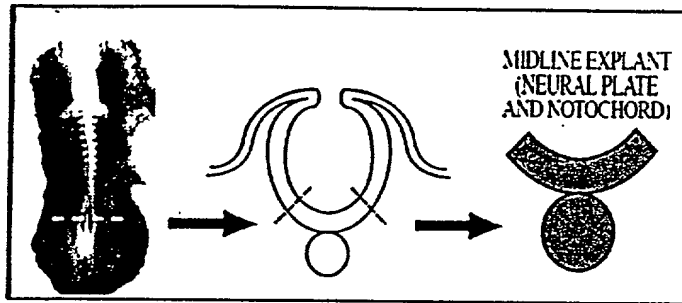


Fig. 3A

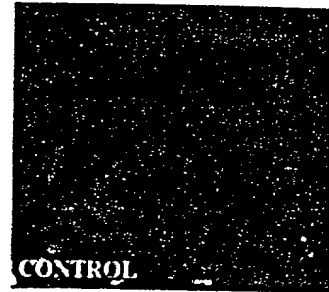


Fig. 3B

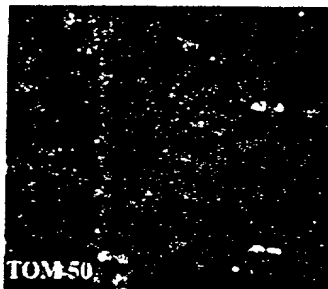


Fig. 3C



Fig. 3D

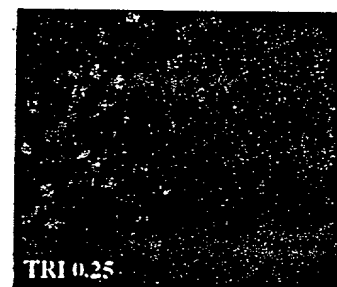


Fig. 3E

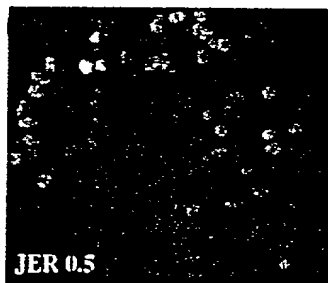


Fig. 3F

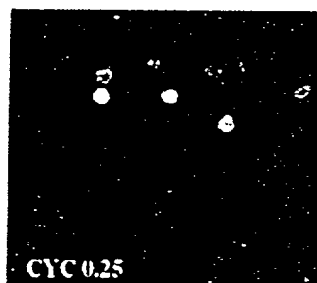


Fig. 3G

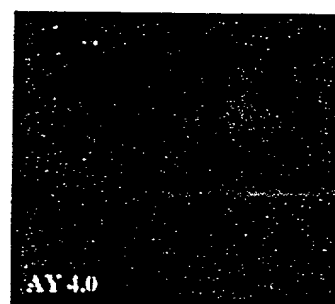


Fig. 3H



Fig. 3I

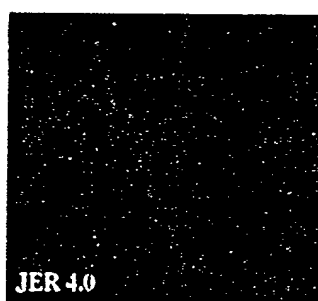


Fig. 3J

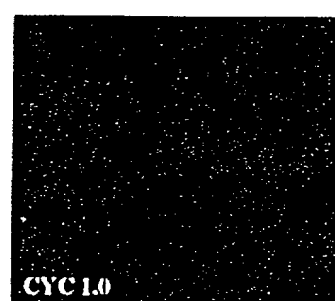


Fig. 3K

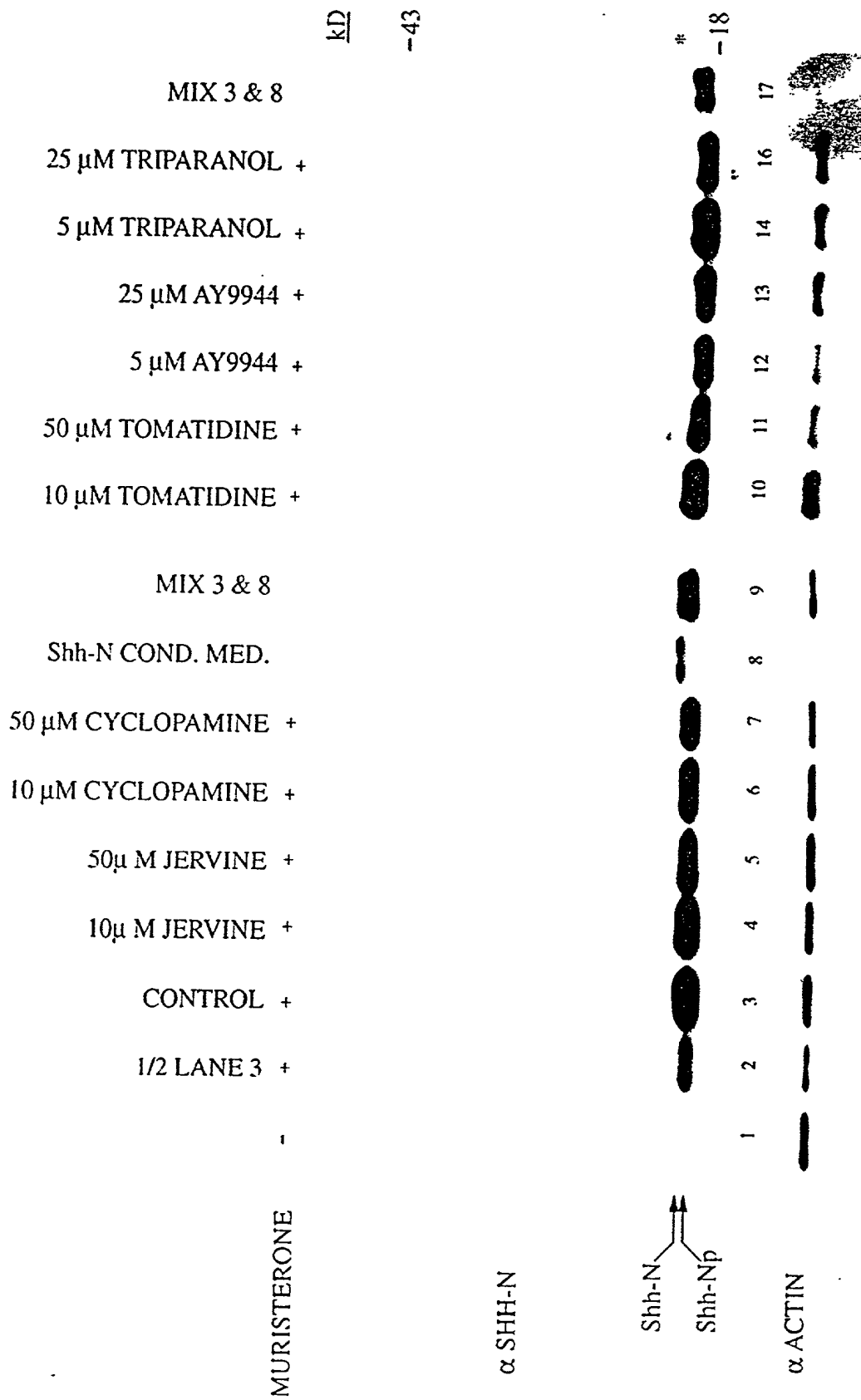


Fig. 4

Fig. 5A

Fig. 5B



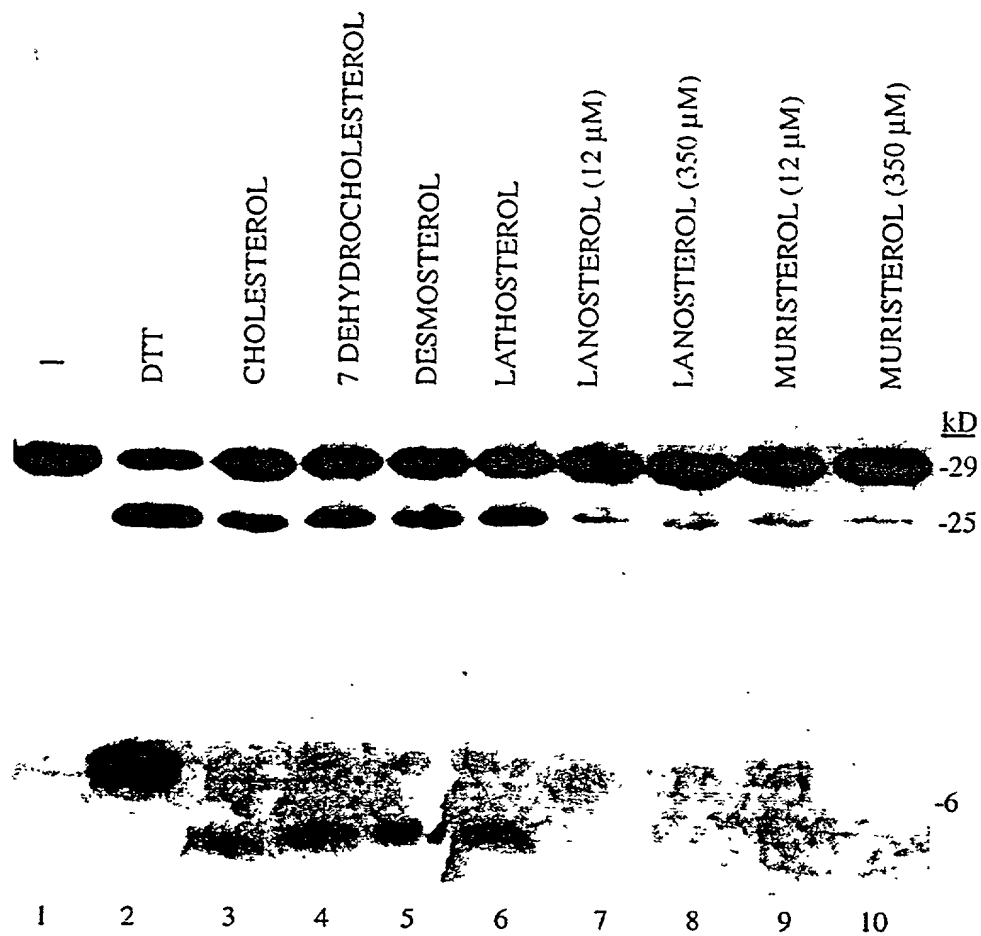


Fig. 5C

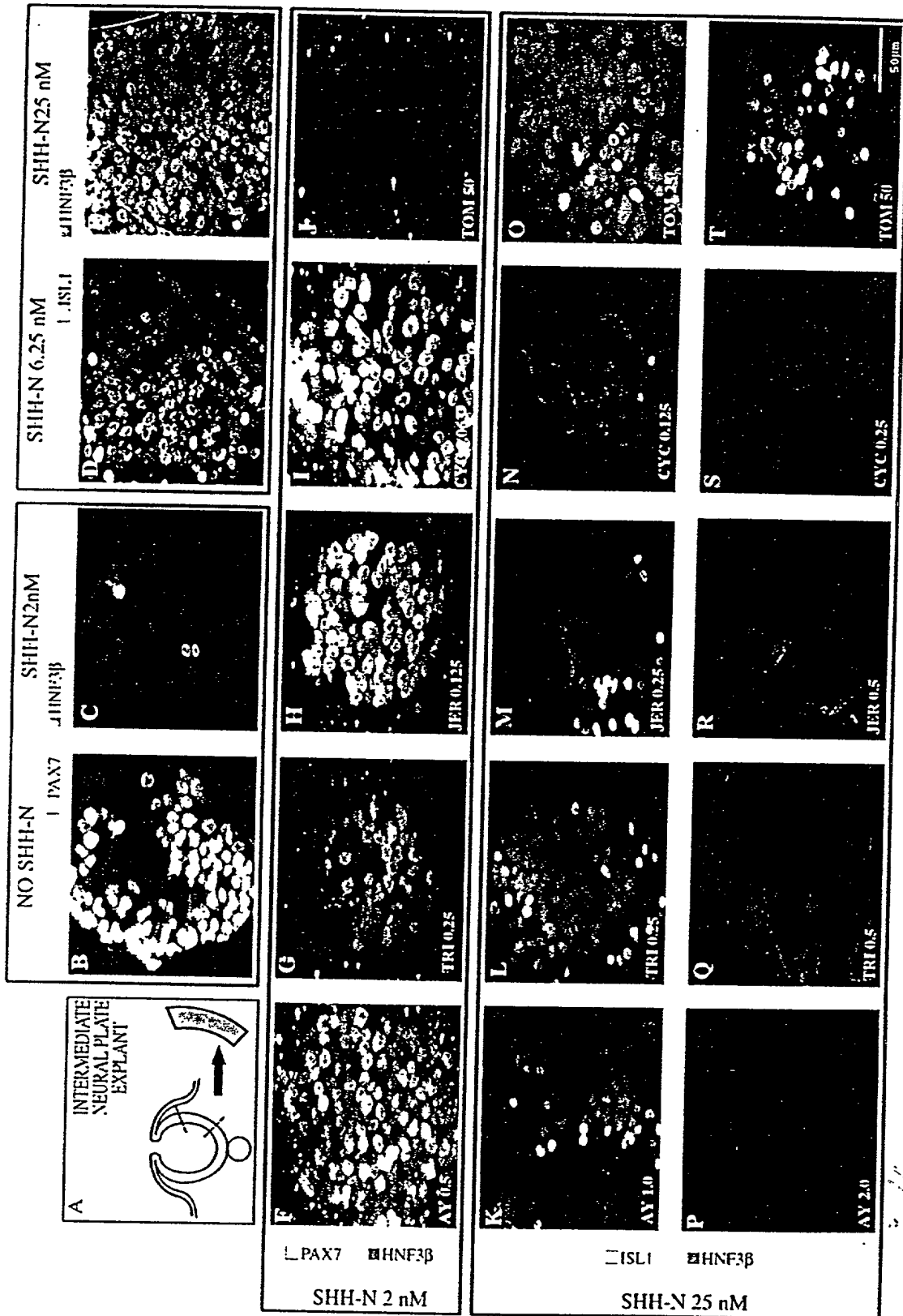


Fig. 6 (A-T)



Fig. 7A

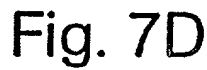
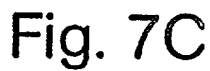
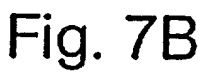


Fig. 8B

*Gli1*

Shh -/-

Fig. 8D



*Ptc1*

Shh -/-

Control

Shh -/-

Fig. 8H

Control

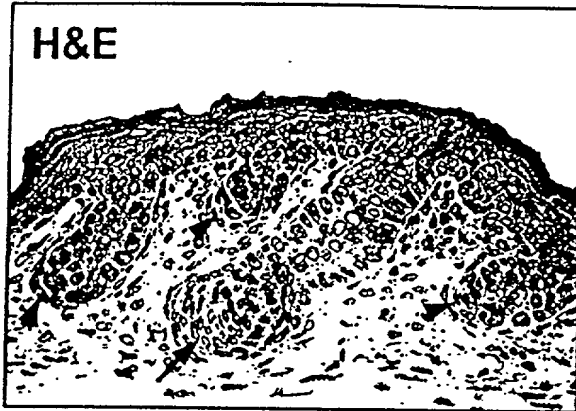


Fig. 9A

*Shh*<sup>-/-</sup>

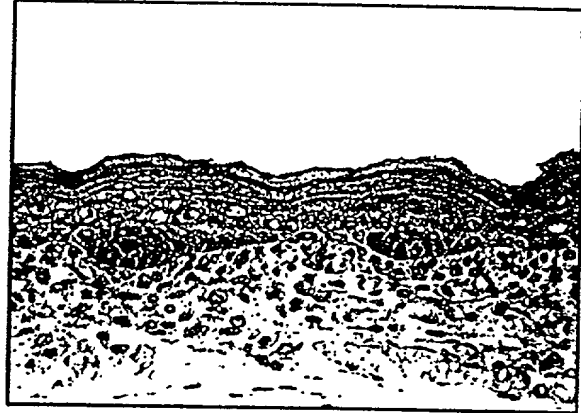


Fig. 9B

Control

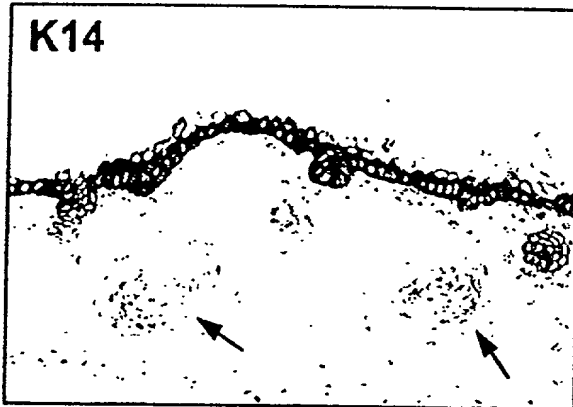


Fig. 9C

*Shh*<sup>-/-</sup>

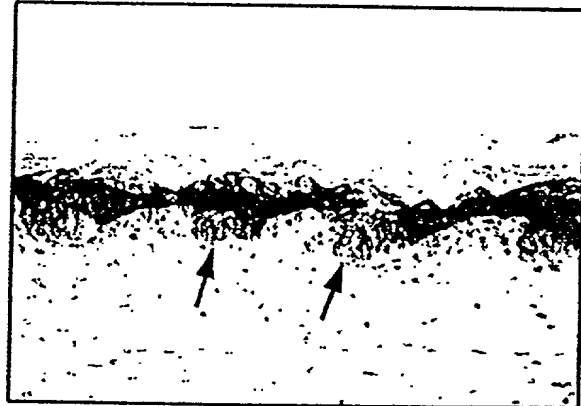


Fig. 9D

Control

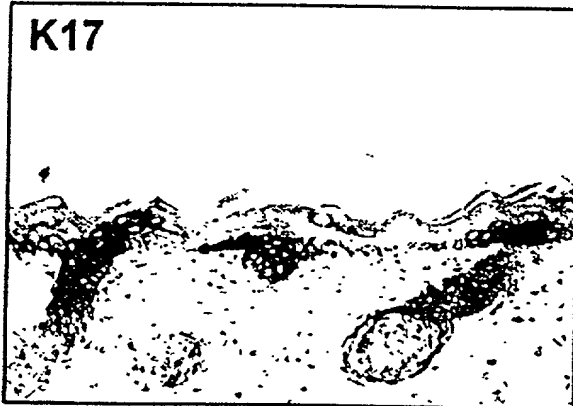


Fig. 9E

*Shh*<sup>-/-</sup>



Fig. 9F

000011-42680260



## Control



Fig. 10B

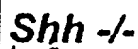


Fig. 10C

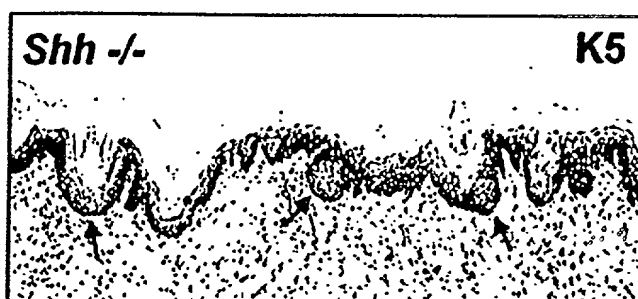


Fig. 10D



Fig. 10E



Fig. 10F

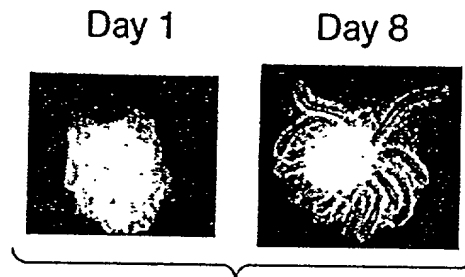


Fig. 11A

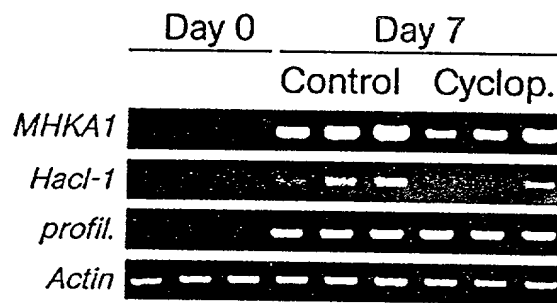
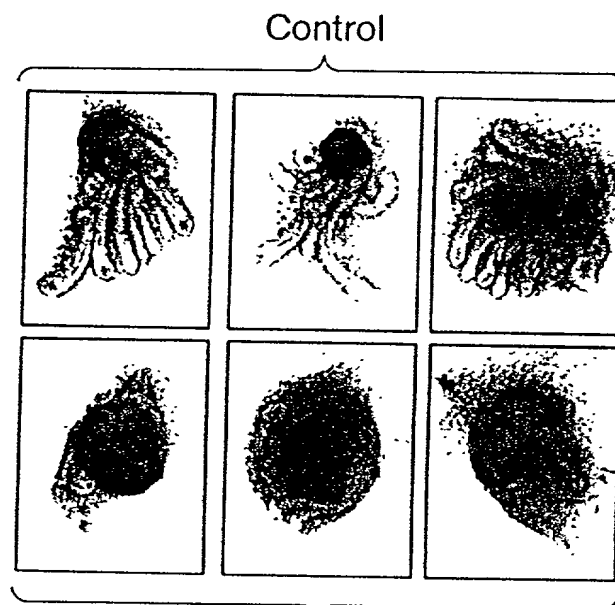


Fig. 11 B



Cyclopamine

Fig. 11C



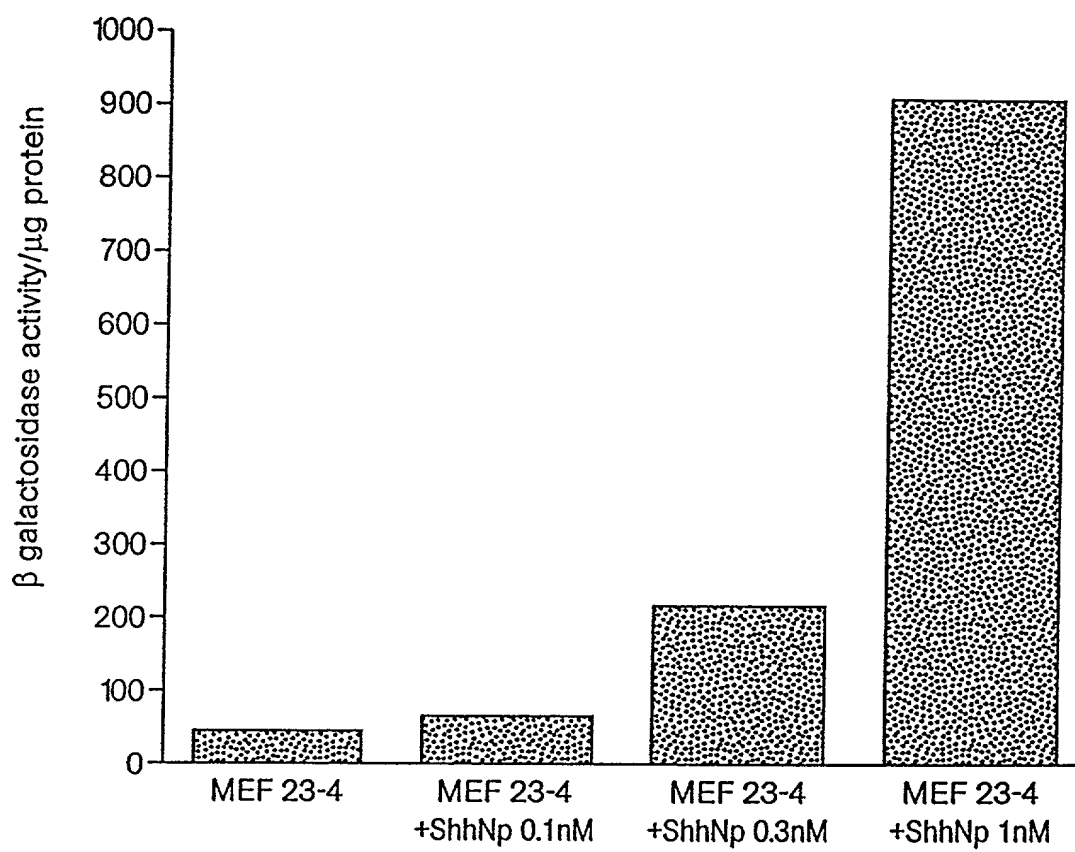


Fig. 12

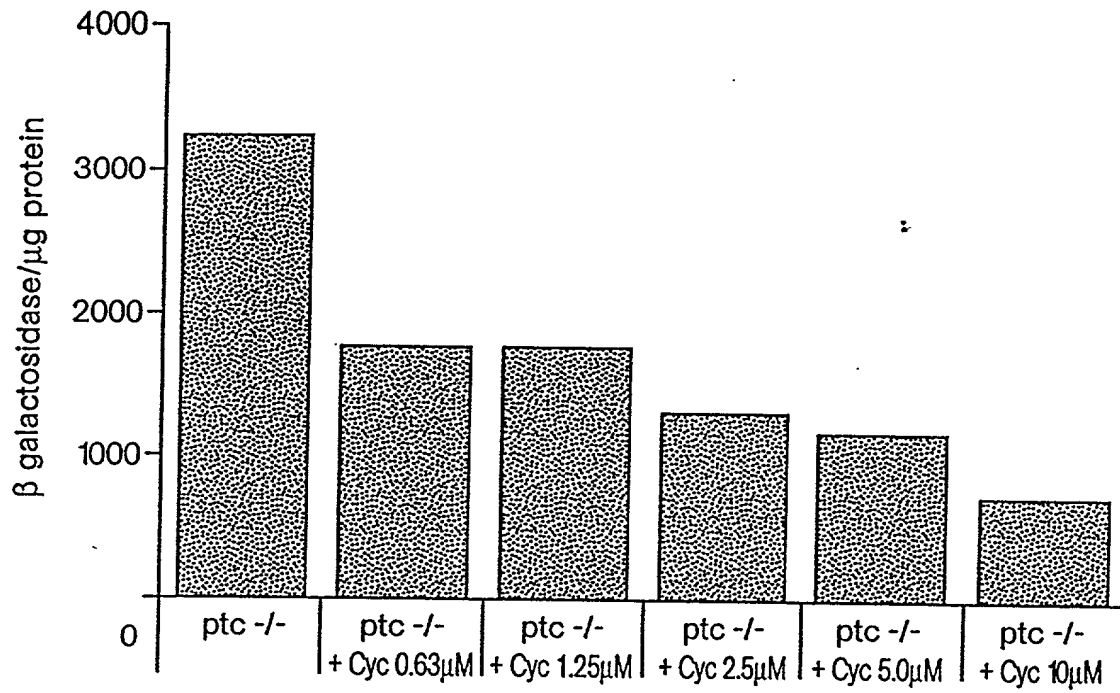


Fig. 13

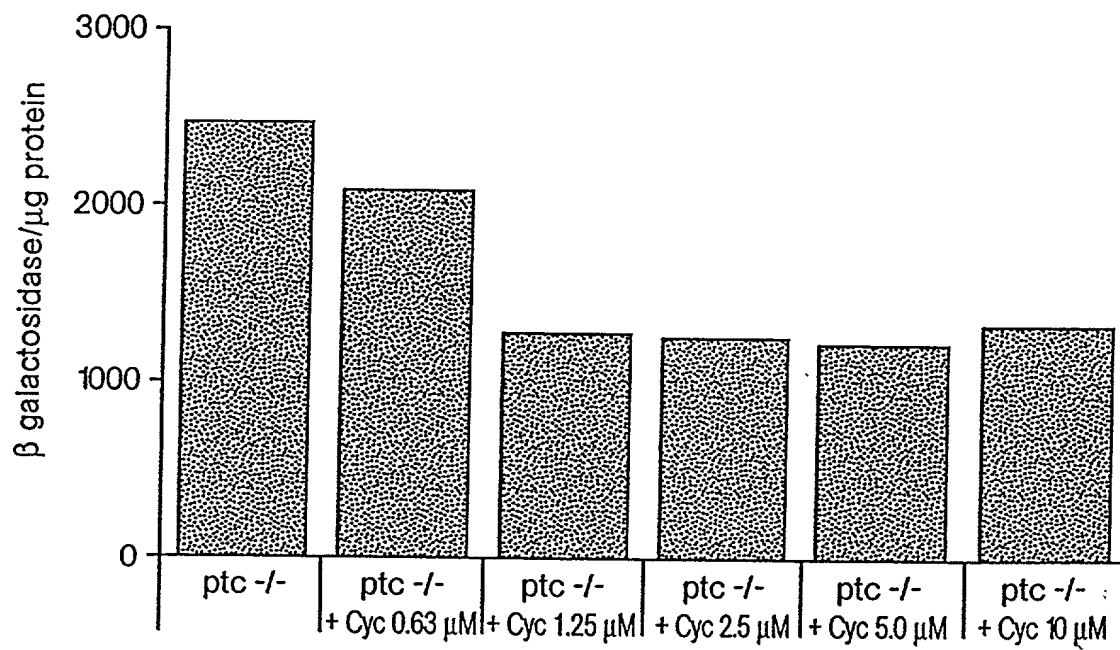


Fig. 14

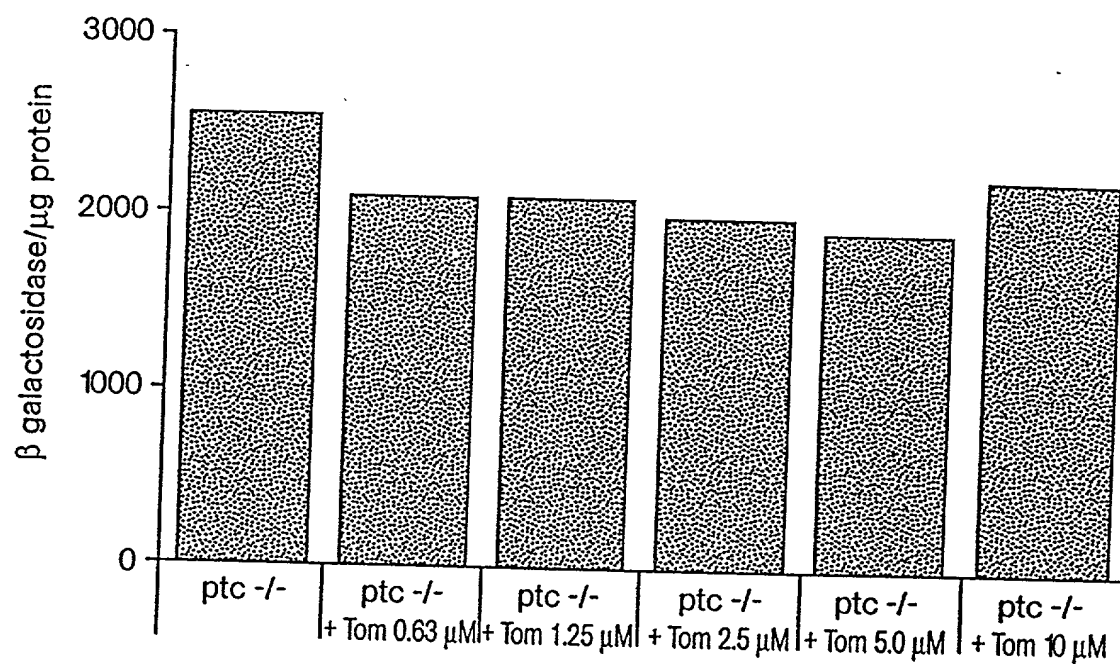


Fig. 15

①



I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

[illegible]

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulation, §1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, § 119(a)-(d) of any foreign application(s) for patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed.

Prior Foreign Application(s)			Priority Claimed
_____	_____	_____	( ) Yes ( ) No
(Number)	(Country)	(Day/Month/Year Filed)	
_____	_____	_____	( ) Yes ( ) No
(Number)	(Country)	(Day/Month/Year Filed)	

I hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States Provisional application(s) listed below.

60/081,186                      9 April 1998  
(Application Number)              (Filing Date)

I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application.

<u>09/090,622</u>	<u>June 4, 1998</u>
(Application Number)	(Filing Date)

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I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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